
Monte Carlo Simulations of the 3-State Potts Model in 2D

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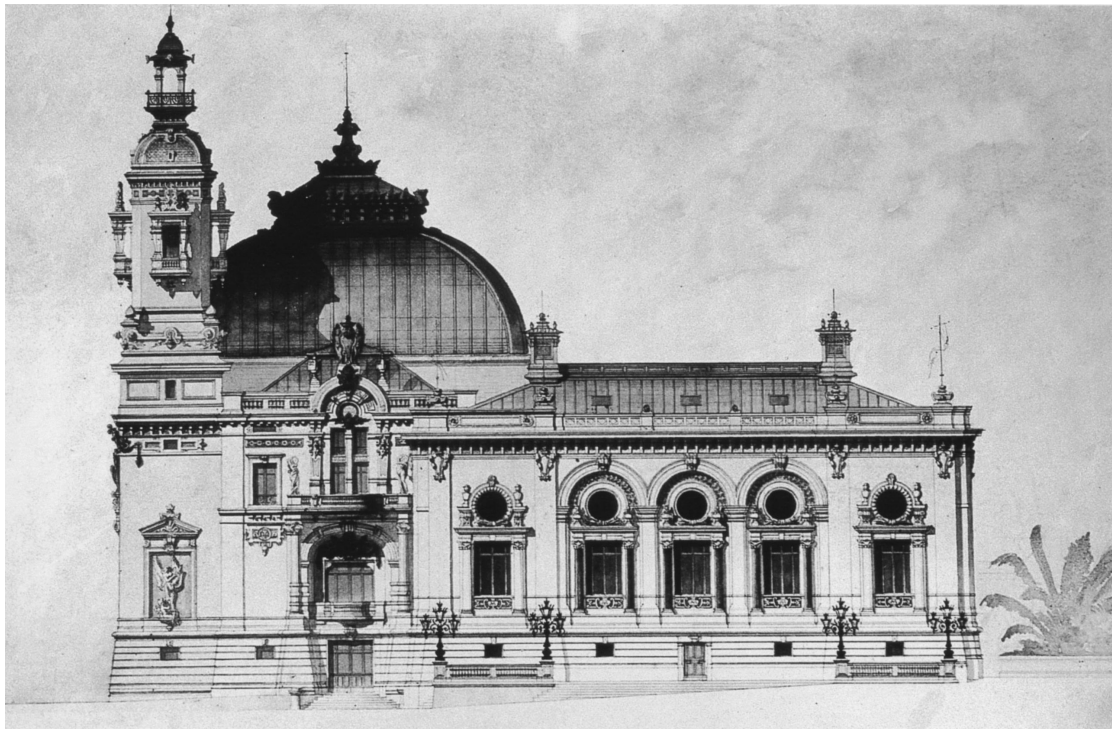
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Image Credit: Eastern frontal view of the Trente-Quarante Gaming Room of the Monte Carlo Casino, from which the methods used in this work get their name, designed by Charles Garnier c. 1878–80. Image retrieved from Wikimedia Commons, originally in Bonillo et al. (2004). *Charles Garnier and Gustave Eiffel on the French and Italian Rivieras: The Dream of Reason*

Abstract

We study the critical behaviour of the 3-state Potts model in two dimensions and zero-field. Monte Carlo simulations are used to find the critical temperature, as well as the critical exponents of the model. These values are calculated by using finite size scaling methods.

The Ising model in two dimensions is used as a benchmark for the methods used, given that it is exactly solved. The critical temperature found corresponds well with the exact known value of Onsager's solution. Values of ratios of critical exponents also agree reasonably well as do critical exponents themselves, when compared to exact results.

For the Potts model the critical temperature is found to be in moderate correspondence with exact results. Ratios of critical exponents agree well with conjectured results. Using the ratio β/ν all critical exponents are in agreement with conjectured results while this is less so using γ/ν , possibly because of finite size scaling corrections. Special attention is paid to scaling for the heat capacity by including a background term.

Populaire Samenvatting

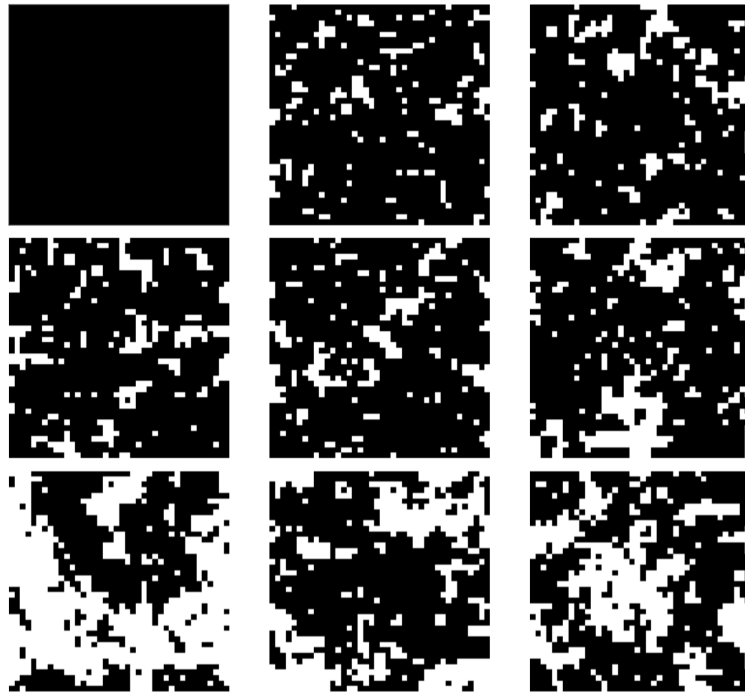
Faseovergangen zien we overal om ons heen. Water verandert in stoom of bevriest tot ijs. Bij een faseovergang zien we dat een bepaald getal, dat we de orde parameter noemen, een sprong maakt. Bij een magneet is dat bijvoorbeeld de magnetisatie, die een waarde heeft bij lage temperaturen maar 0 wordt als de magneet wordt opgewarmd tot boven zijn Curie temperatuur.

Faseovergangen worden getypeerd door de temperatuur waarop ze plaatsvinden, de zogenaamde kritische temperatuur, en door kritische exponenten. Dit zijn getallen die bepalen hoe snel een eigenschap van een systeem verandert dicht bij de faseovergang.

In dit verslag beschrijf ik de faseovergang van volledig gemagnetiseerde naar ongemagnetiseerde toestand van twee modellen en hoe ik heb geprobeerd hun kritische temperatuur en exponenten te vinden. Het Ising model beschrijft een rooster waarbij op ieder roosterpunt een klein magneetje zit (dat we een spin noemen), dat omhoog of omlaag kan wijzen. Het Potts model veralgemeniseert dit idee door iedere spin in een richting te laten wijzen, waarbij de mogelijke richtingen gelijk verdeelt zijn over een cirkel. Ik heb specifiek gekeken naar een Potts model met drie mogelijke richtingen.

Om de faseovergangen te onderzoeken heb ik twee verschillende zogenaamde Monte Carlo algoritmen gebruikt om de systemen te simuleren. De eerste was het Metropolis algoritme (In het plaatje zie je hoe het rooster verandert als het met dit algoritme gesimuleerd wordt.) Hierbij wordt voor de spins op het rooster een voor een geprobeerd ze om te draaien. Dit gebeurt met een bepaalde kans. Omdat het in de buurt van de faseovergang kan gebeuren dat er grote eilanden ontstaan van spins die allemaal dezelfde richting op wijzen en waarvan de kans klein is dat ze worden omgedraaid, heb ik ook gebruik gemaakt van het Wolff algoritme dat grote clusters van spins in één keer omdraait. Zo kan het niet gebeuren dat het rooster tijdens de simulatie lange tijd in dezelfde toestand blijft.

Tijdens het draaien van de simulaties deed ik continue metingen van de energie en magnetisatie. Met behulp van die metingen kon ik verschillende grootheden uitrekenen zoals de warmtecapaciteit. Om de data te analyseren en de kritische temperatuur en exponenten te bepalen heb ik gebruik gemaakt van een eigenschap die dit soort systemen hebben. Als je roosters van verschillende grootten simuleert en dan de waarde van



bepaalde eigenschappen met een factor (die afhangt van de grootte van het rooster, de kritische temperatuur en de kritische exponenten) vermenigvuldigt, blijkt dat de verschillende lijnen van verschillende rooster grootten precies over elkaar heen vallen. Dit gebeurt wel alleen als je de factor met de juiste exponenten en temperatuur berekent. Door verschillende waarden te proberen kun je zo de exponenten vinden, door te kijken wanneer de metingen van verschillende roostergrootten het best samenvallen.

Voor zowel het Ising model als het Potts model heb ik de kritische temperatuur kunnen vinden. Ook kon ik voor beiden nauwkeurig de verhouding tussen kritische exponenten bepalen. Bij het Ising model vond ik kritische exponenten die goed overeenkomen met de exacte waarden. Voor het Potts model vond ik, met één manier om data samen te laten vallen, exponenten die overeenkomen met voorspelde waarden. De andere manier gaf juist resultaten die niet goed waren. Dit komt mogelijk doordat de systemen die ik simuleerde relatief klein zijn, terwijl deze technieken beter werken voor grote systemen.

Contents

Introduction	5
1 Models and Critical Phenomena	6
1.1 Phase Transitions and Critical Phenomena	6
1.1.1 Correlation Functions	7
1.1.2 Scaling Laws	8
1.1.3 Universality	9
1.2 The Two-Dimensional Ising Model	9
1.2.1 Thermodynamic Properties of the Two-Dimensional Ising Model .	11
1.2.2 The Ising Model around T_c	13
1.2.3 Magnetisation of the Ising Model	14
1.3 The Potts Model	15
2 Simulating Lattice Models	17
2.1 Markov Processes and Monte Carlo Methods	17
2.2 The Metropolis Algorithm	19
2.2.1 Optimisations	21
2.2.2 Critical Slowing Down	22
2.3 The Wolff Algorithm	22
2.3.1 Generalising Algorithms to the Potts Model	23
3 Analysing Monte Carlo Simulations	25
3.1 Statistics	25
3.1.1 Binning	25
3.1.2 Bootstrap and Jackknife	26
3.2 Finite Size Scaling	27
3.3 The Binder Cumulant	28
4 Results	32
4.1 The Ising Model at Criticality	32
4.2 The Potts Model at Criticality	34
5 Conclusions	39
5.1 Further Work	39
5.2 Acknowledgements	40
Bibliography	40

Introduction

Phase transitions are an everyday part of life, the most well known being the transition of water into water vapour or ice into water. They also occur in more abstract contexts such as networks of neurons [1].

Although there are some systems that can be exactly solved and for which all properties are known, including at the phase transition, many more cannot, and thus different methods have to be employed to understand these transitions. In this thesis we will use Monte Carlo simulations to study the critical behaviour of the 3-state Potts model. Specifically we are interested in the critical exponents and critical temperature of the model. We proceed as follows:

In Chapter 1 we consider exactly what properties a phase transition has. We see that the behaviour of many thermodynamic properties of a system at criticality can be described by critical exponents. These exponents are closely related to each other. We also consider the two-dimensional Ising model in zero-field, which has been solved exactly and for which the critical exponents are known. Then we have a look at the Potts model, its definition and some conjectures concerning properties of the phase transition.

In Chapter 2 we look at the theory behind simulating lattice systems using Monte Carlo simulations. We define the Metropolis algorithm, what shortcomings it has and how those shortcomings can be amended by using the Wolff algorithm. We also consider some optimisations for the simulations.

Chapter 3 lays out the way to analyse the data obtained from the simulation. Some care has to be taken to use statistically independent measurements. We also look at resampling methods used to calculate errors of properties of the system that depend in a complicated way on the measured quantities. The critical temperature is found by using Binder cumulants, while critical exponents are found using finite size scaling methods.

In Chapter 4 the results of applying the methods from Chapter 3 are presented for the two-dimensional Ising and Potts models. The measured values for the critical temperatures and exponents agree reasonably well with the exact solution (for the Ising model) and other simulation and conjectures (for the Potts model).

In Chapter 5 we summarise the results and give an outlook for further work. Due to the strong computational component of this project, a significant chunk of code is not included here. A repository containing the code used, the data sets that were generated and a short usage manual may be found at github.com/teunzward/bachelor-project.

Chapter 1

Models and Critical Phenomena

1.1 Phase Transitions and Critical Phenomena

To determine when a phase transition has taken place, we consider the order parameter. In ferromagnetic systems such as iron, as well as in the systems we will study in this work, this is the magnetisation. On one side of the phase transition a non-zero magnetisation is present. As the iron is heated, it moves past its Curie temperature and the magnetisation does become zero. More generally we consider an order parameter ϕ which is a quantity that is non-zero on one side of the phase transition, where a symmetry of the Hamiltonian governing this system is spontaneously broken, and vanishes on the other side. Usually the order parameter is zero on the high temperature side of the phase transition. (See figure 1.1. The Ising model is considered in more detail in section 1.2.)

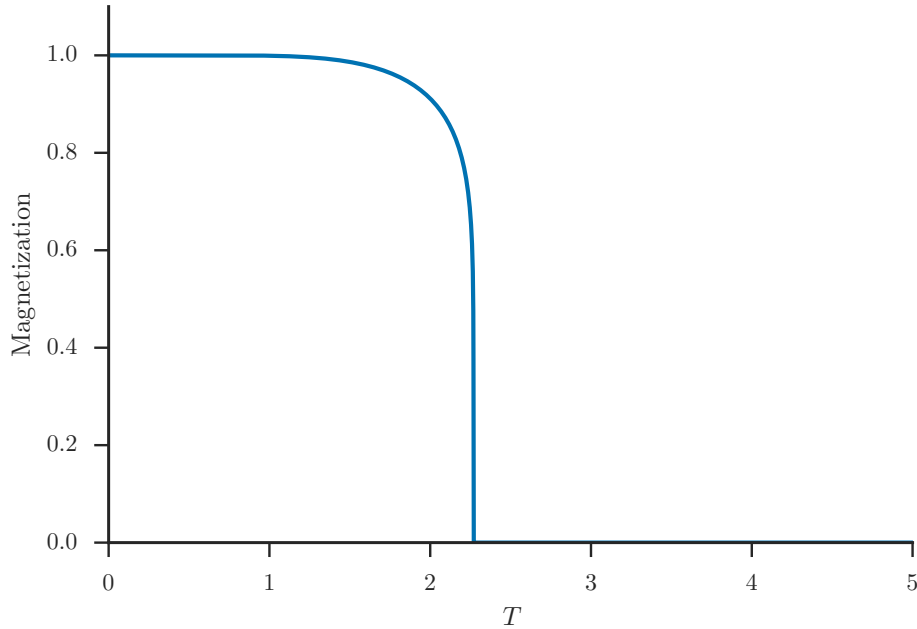


Figure 1.1: The magnetisation of the two-dimensional Ising model as a function of temperature. The critical temperature is 2.269. Notice how the magnetisation is finite on one side of the phase transition, but zero on the other side.

When we consider phase transitions, we distinguish two different kinds. The phase

transition associated with freezing water is what is called first-order. When the critical temperature is crossed the water molecules move from a disordered phase into an ordered crystal phase. As this happens, energy is emitted in the form of latent heat. This is defined as

$$l = \int_{T_c^-}^{T_c^+} c(T) dT, \quad (1.1)$$

with l the latent heat and $c(T)$ the heat capacity of the system. In first order transitions the order parameter is discontinuous at T_c [2]. In the rest of this work we only consider second-order transitions, for which the latent heat is zero and the order parameter, but not the rate of change of the order parameter, is continuous at T_c .

A phase transition is usually also characterised by the behaviour of thermodynamic properties of the system, such as the specific heat or the magnetic susceptibility. Often the heat capacity diverges as $c \propto |T - T_c|^{-\alpha}$. We call α a critical exponent. Because for continuous phase transitions the latent heat has to vanish by definition, α has to be smaller than 1, because otherwise the integral of equation (1.1) diverges. No divergence occurs if $\alpha < 0$. In the limiting case that $\alpha = 0$, we can consider the divergence of the specific heat to be logarithmic since

$$\log\left(\frac{1}{x}\right) = \lim_{\alpha \rightarrow 0^+} \frac{1}{\alpha} (x^{-\alpha} - 1), \quad (1.2)$$

where $x = |T - T_c|/T_c$. Theoretically and experimentally we find that the exponent governing divergence is the same both above and below T_c [2].

1.1.1 Correlation Functions

For the systems we would like to consider it is often interesting to quantify how different parts of system relate to each other. In fact we will see that this correlation proves critical when choosing an appropriate algorithm to numerically study systems around the critical temperature (section 2.2.2). To quantify the correlations in a system we define the two-point correlation function [2], which, to illustrate some properties of the correlation function, we first define for a system of spins:

$$G^{(2)}(\mathbf{i}, \mathbf{j}) = \langle \mathbf{s}_i \cdot \mathbf{s}_j \rangle. \quad (1.3)$$

Here \mathbf{i} and \mathbf{j} are the position vectors of the spins at locations i and j respectively. The angle brackets denote thermal averaging. Because the system is often translationally invariant as well as isotropic, meaning it has no preferred direction, such as in a crystal lattice or in disordered systems, $G^{(2)}$ often depends only on $|\mathbf{i} - \mathbf{j}|$.

In the general case the two-point correlation function is defined as

$$G^{(2)}(r) = \langle \phi(0) \cdot \phi(\mathbf{r}) \rangle, \quad (1.4)$$

where ϕ is the order parameter of the system. Below T_c the system is ordered and $G^{(2)}$ is large for all r . In the example using spins given above, this means almost all spins are aligned. A more useful quantity in this case is the connected correlation function

$$G_c^{(2)} = \langle \phi(0) \cdot \phi(\mathbf{r}) \rangle - |\langle \phi \rangle|^2. \quad (1.5)$$

By subtracting the thermally averaged value of the order parameter from the two-point correlation function, we can ignore the general alignment of the order parameter and only have fluctuations in the order parameter contribute.

When T/T_c is either large or small $G_c^{(2)}$ is small. Precisely at the critical temperature $G_c^{(2)}$ has the form

$$G_c^{(2)} \propto \frac{1}{r^{d-2+\eta}}, \quad (1.6)$$

with d the dimensionality of the system and η another critical exponent. Far away from T_c $G_c^{(2)}$ decays exponentially, but for $|T - T_c|/T_c \ll 1$ $G_c^{(2)}$ has the form

$$G_c^{(2)} \propto \frac{e^{-r/\xi}}{r^{d-2+\eta}}. \quad (1.7)$$

Here ξ denotes the correlation length. Fluctuations of the order parameter up to this length scale are common, but larger fluctuations are exponentially suppressed. The correlation length diverges as T_c is approached from above or below, according to

$$\xi \propto |T - T_c|^{-\nu}, \quad (1.8)$$

where ν is another critical exponent.

1.1.2 Scaling Laws

We can define three other critical exponents. For the magnetic susceptibility we have

$$\chi \propto |T - T_c|^{-\gamma}, \quad (1.9)$$

while for the magnetisation of a system we have

$$m \propto |T - T_c|^\beta \quad \text{as } T \rightarrow T_c \text{ from below.} \quad (1.10)$$

Finally we can define a critical exponent for the magnetisation in a non-zero magnetic field (the previous five critical exponents assumed $B = 0$). Precisely at T_c

$$m \propto B^{1/\delta} \quad \text{as } B \rightarrow 0. \quad (1.11)$$

These six critical exponents are not independent of each other but are related through scaling laws. Given these scaling laws usually only two critical exponents need to be known to determine all others as well. They are:

$$\alpha + 2\beta + \gamma = 2 \quad (1.12)$$

$$\alpha + \beta(\delta + 1) = 2 \quad (1.13)$$

$$(2 - \eta)\nu = \gamma \quad (1.14)$$

$$\nu d = 2 - \alpha, \quad (1.15)$$

with d the dimensionality of the system [2–4]. In the case where only α and ν are known, no other exponent can be determined using these relations, because then we have a system of three equations and four unknowns (the last scaling law only contributes a tautology).

1.1.3 Universality

One property of critical phenomena that simplifies their study is the concept of universality. This is the independence of many thermodynamic properties of a system on the exact details of the Hamiltonian of that system. They will only depend on the dimensionality of the system and symmetries of the Hamiltonian. Consider the Hamiltonian

$$H(s) = H_0(s) + \lambda H_1(s), \quad (1.16)$$

where s denotes the state of the system, H_0 is a part with a given symmetry, and H_1 does not have that symmetry. The critical exponents then only depend on λ in that they have one value when $\lambda = 0$ and another when $\lambda \neq 0$. This also means that, if both H_0 and H_1 have the same symmetries, then, if H_0 is some simple Hamiltonian while H_1 is more complicated, it is possible to obtain the critical exponents of a system using the simple part while stripping the more complicated part. This may significantly ease the determination of critical exponents when performing simulations. Different systems with the same critical exponents are said to be in the same universality class. It is believed (and experimentally established within error) that such diverse systems as carbon dioxide, xenon and the three-dimensional Ising model are in the same universality class [3].

1.2 The Two-Dimensional Ising Model

To validate methods to determine critical exponents for systems that are not exactly solvable, a control system that is exactly solved is useful. To that end we study the two-dimensional Ising model in zero-field, first solved exactly in 1944 by Lars Onsager [5]. It describes a square lattice with nearest neighbour interactions, where each lattice point has with it associated a number (which we will refer to as spin) which may either be $+1$ or -1 and was originally meant as a model for magnets.¹ The Hamiltonian in zero-field is [6]

$$H = -J_1 \sum_{j=1}^{\mathcal{M}} \sum_{k=1}^{\mathcal{N}} \sigma_{j,k} \sigma_{j,k+1} - J_2 \sum_{j=1}^{\mathcal{M}} \sum_{k=1}^{\mathcal{N}} \sigma_{j,k} \sigma_{j+1,k}, \quad (1.17)$$

with \mathcal{M} and \mathcal{N} the extent of the lattice in the x - and y -directions respectively and J_1 and J_2 the interaction strength between neighbours in respectively the x - and y -directions. In the case where the interaction strength in both directions is the same the Hamiltonian becomes [7]

$$H = -J \sum_{\langle ij \rangle} \sigma_i \sigma_j. \quad (1.18)$$

The sum runs over all nearest neighbour bonds on the lattice. This Hamiltonian is symmetric under a change of sign of all spins. In the ferromagnetic ground state ($J > 0$, $T = 0$) all spins on the lattice are aligned in one of the two possible directions (the direction is chosen when the mirror symmetry in the lattice plane is spontaneously broken as the lattice cools). The Hamiltonian is subject to periodic boundary conditions in both directions, meaning $\sigma_{1,k} = \sigma_{\mathcal{M}+1,k}$ and $\sigma_{j,1} = \sigma_{j,\mathcal{N}+1}$. We are interested in

¹The model is a better description of adsorption of gases on the surface of crystals [5].

the thermodynamic properties of the Ising Model. To that end we define the partition function

$$Z = \sum_{\sigma=\pm 1} e^{-\beta H} \quad (1.19)$$

$$= \sum_{\sigma=\pm 1} \prod_{j=1}^{\mathcal{M}} \prod_{k=1}^{\mathcal{N}} e^{\beta J_1 \sigma_{j,k} \sigma_{j,k+1}} \prod_{j=1}^{\mathcal{M}} \prod_{k=1}^{\mathcal{N}} e^{\beta J_2 \sigma_{j,k} \sigma_{j+1,k}}, \quad (1.20)$$

with $\beta = 1/k_B T$ and the sum running over every possible orientation of the spins on the lattice. Solving this in the thermodynamic limit requires a non-trivial amount of effort and it is best to refer to either Onsager [5], who systemically added one-dimensional Ising models together to create a two dimensional lattice, or Kasteleyn as described in [6], whose approach considerably simplifies the derivation by reducing it to a combinatorial problem. For small systems (for example a 3 by 3 lattice) the sum could be explicitly calculated, but the number of terms grows exponentially as 2^N [2, 7].

The derivation introduces a sign ambiguity in Z which takes some additional care to resolve, but we can avoid having to deal with this by considering the free energy F in the thermodynamic limit² instead of the partition function

$$\begin{aligned} F &= -\frac{1}{\beta} \lim_{\substack{\mathcal{N} \rightarrow \infty \\ \mathcal{M} \rightarrow \infty}} \frac{1}{\mathcal{M}\mathcal{N}} \log(Z_{\mathcal{M},\mathcal{N}}) \\ &= -\frac{1}{\beta} \left[\log(2) + \frac{1}{2} \frac{1}{(2\pi)^2} \int_0^{2\pi} d\theta_1 \int_0^{2\pi} d\theta_2 \log \left(\cosh(2\beta J_1) \cosh(2\beta J_2) \right. \right. \\ &\quad \left. \left. - \sinh(2\beta J_1) \cos(\theta_1) - \sinh(2\beta J_2) \cos(\theta_2) \right) \right]. \end{aligned} \quad (1.21)$$

F is an analytic function of the temperature T , except at one value, which we will call the critical temperature T_c . At this temperature we can define the equality [6]

$$|z_1| = \frac{1 - |z_2|}{1 + |z_2|}, \text{ with } z_1 = \tanh(2\beta J_1), z_2 = \tanh(2\beta J_2). \quad (1.22)$$

Rewriting and squaring this gives

$$1 - |z_1 z_2| = |z_1| + |z_2| \rightarrow \quad (1.23)$$

$$(1 - z_1^2)(1 - z_2^2) = 4|z_1 z_2| \quad (1.24)$$

Finally, using

$$\frac{1}{2z_k} (1 - z_k^2) = \frac{1}{\sinh(2\beta J_k)}, \text{ with } k \in \{1, 2\} \quad (1.25)$$

we get the equality (at the critical point)

$$1 = \sinh(2\beta J_1) \sinh(2\beta J_2). \quad (1.26)$$

²This is the limit in which the number of particles on the lattice tends to infinity.

In the case where interaction strength in both the x - and y -directions is the same ($|J_1| = |J_2| = J$) we get an expression for the critical temperature in terms of the bond energy

$$1 = \sinh(2\beta J) \rightarrow \quad (1.27)$$

$$k_B T_c = \frac{2}{\operatorname{asinh}(1)} J = \frac{2}{\log(1 + \sqrt{2})} J \approx 2.269 J \quad (1.28)$$

Onsager [5] also calculated the values of the free energy, internal energy and entropy at the critical temperature³:

$$-\frac{f_c}{k_B T} = \frac{1}{2} \log 2 + \frac{2}{\pi} G \approx 0.929, \quad (1.29)$$

$$u_c = -\sqrt{2} J \approx -1.414 J, \quad (1.30)$$

$$\frac{s_c}{k_B} = \log(\sqrt{2} e^{2G/\pi}) - \sqrt{2} \frac{J}{k_B T} \approx 0.306 \quad (1.31)$$

with G Catalan's constant.⁴

1.2.1 Thermodynamic Properties of the Two-Dimensional Ising Model

From the free energy it is relatively simple to determine the specific heat per spin and the internal energy per spin by taking the appropriate derivatives of the free energy [6]. We work with the isotropic case where $|J_1| = |J_2| = J$ and $z_1 = z_2 = z = \tanh(\beta J)$, corresponding to the Hamiltonian of equation (1.18). Then the expression for the free energy is

$$F = -\frac{1}{\beta} \left[\log(2 \cosh^2(\beta J)) + \log(1 + z^2) + \frac{1}{2\pi^2} \int_0^\pi d\theta_1 \int_0^\pi d\theta_2 \log\left(1 - \frac{1}{2}k \{\cos(\theta_1) + \cos(\theta_2)\}\right) \right], \quad (1.32)$$

with

$$k = \frac{4z(1 - z^2)}{(1 + z^2)^2} = \frac{2 \sinh(2\beta J)}{\cosh^2(2\beta J)}. \quad (1.33)$$

Performing the substitutions $\omega_1 = \frac{1}{2}(\theta_1 + \theta_2)$ and $\omega_2 = \frac{1}{2}(\theta_1 - \theta_2)$ and integrating over ω_1 the free energy becomes

$$F = -\frac{1}{\beta} \left[\log(2 \cosh(2\beta J)) + \frac{1}{\pi^2} \int_0^{\frac{\pi}{2}} d\omega_1 \int_0^{\frac{\pi}{2}} d\omega_2 \log(1 - k \cos(\omega_1) \cos(\omega_2)) \right] \quad (1.34)$$

$$= -\frac{1}{\beta} \left[\log(\sqrt{2} \cosh(2\beta J)) + \frac{1}{\pi} \int_0^{\frac{\pi}{2}} d\omega \log\left(1 + \left\{1 - k^2 \cos^2(\omega)\right\}^{\frac{1}{2}}\right) \right] \quad (1.35)$$

³We use lowercase letters to denote the thermodynamic properties of a single spin on the lattice and uppercase letters when referring to the entire lattice. Taking as an example the internal energy, $U/N = u$ with N the number of spins on the lattice.

⁴ $G = 1^{-2} - 3^{-2} + 5^{-2} - 7^{-2} + \dots \approx 0.916$

We will take derivatives from this expression to determine expressions for thermodynamic properties. The internal energy per spin is

$$u = \frac{\partial \beta F}{\partial \beta} \quad (1.36)$$

$$= -2J \tanh(2\beta J) + k \frac{dk}{d\beta} \frac{1}{\pi} \int_0^{\frac{\pi}{2}} d\omega \frac{\sin^2(\omega)}{\Delta(1+\Delta)} \quad (1.37)$$

with

$$\Delta = \left(1 - k^2 \sin^2(\omega)\right)^{\frac{1}{2}}. \quad (1.38)$$

Note that the argument of the integral in equation (1.36) can be rewritten as

$$\frac{\sin^2(\omega)}{\Delta(1+\Delta)} = \frac{(1-\Delta)\sin^2(\omega)}{\Delta(1-\Delta^2)} = \frac{1}{k^2} \left(\frac{1}{\Delta} - 1\right), \quad (1.39)$$

from which it follows that

$$u = -2J \tanh(2\beta J) + \frac{1}{\pi} \frac{1}{k} \frac{dk}{d\beta} \left[K(k) - \frac{\pi}{2} \right]. \quad (1.40)$$

Here $K(k)$ is the complete elliptic integral of the first kind

$$K(k) = \int_0^{\frac{\pi}{2}} \frac{d\phi}{\left(1 - k^2 \sin^2(\phi)\right)^{\frac{1}{2}}}. \quad (1.41)$$

Taking the derivative of k

$$\frac{1}{k} \frac{dk}{d\beta} = \frac{2J}{\tanh(2\beta J)} (1 - 2 \tanh^2(2\beta J)), \quad (1.42)$$

the internal energy per spin becomes (see figure 1.2 for a plot)

$$u = \frac{-J}{\tanh(2\beta J)} \left[1 + \frac{2}{\pi} \left\{ 2 \tanh^2(2\beta J) - 1 \right\} K(k) \right]. \quad (1.43)$$

For the specific heat per spin we take the derivative of the internal energy per spin with respect to the temperature

$$c = \frac{\partial u}{\partial T} = \frac{-1}{k_B T^2} \frac{\partial u}{\partial \beta} \quad (1.44a)$$

$$\begin{aligned} &= \frac{J}{k_B T^2} \left[\frac{-2J}{\cosh^2(2\beta J)} \left\{ 1 + \frac{2}{\pi} \left(2 \tanh^2(2\beta J) - 1 \right) K(k) \right\} \right] \\ &+ \frac{16}{\pi} \frac{J}{\tanh^2 2\beta J} K(k) + \frac{2}{\pi} \frac{\left(2 \tanh^2(2\beta J) - 1 \right)}{\tanh(2\beta J)} \frac{dk}{d\beta} \frac{dK(k)}{dk}. \end{aligned} \quad (1.44b)$$

For the derivative of the elliptic integral we use the identity [6]

$$\frac{dK(k)}{dk} = \frac{1}{k k'^2} \left[E(k) - k'^2 K(k) \right], \quad (1.45)$$

where $E(k)$ is the complete elliptic integral of the second kind

$$E(k) = \int_0^{\frac{\pi}{2}} d\phi \left(1 - k^2 \sin^2(\phi)\right)^{\frac{1}{2}}, \quad (1.46)$$

and $k'^2 = 1 - k^2$. Using equation (1.42) and equation (1.46) the expression for the specific heat per spin becomes (see figure 1.2 for a plot)

$$c = k_B \left[\frac{\beta J}{\tanh(2\beta J)} \right]^2 \frac{2}{\pi} \left[2K(k) - 2E(k) - (1 - k') \left\{ \frac{\pi}{2} + k'K(k) \right\} \right]. \quad (1.47)$$

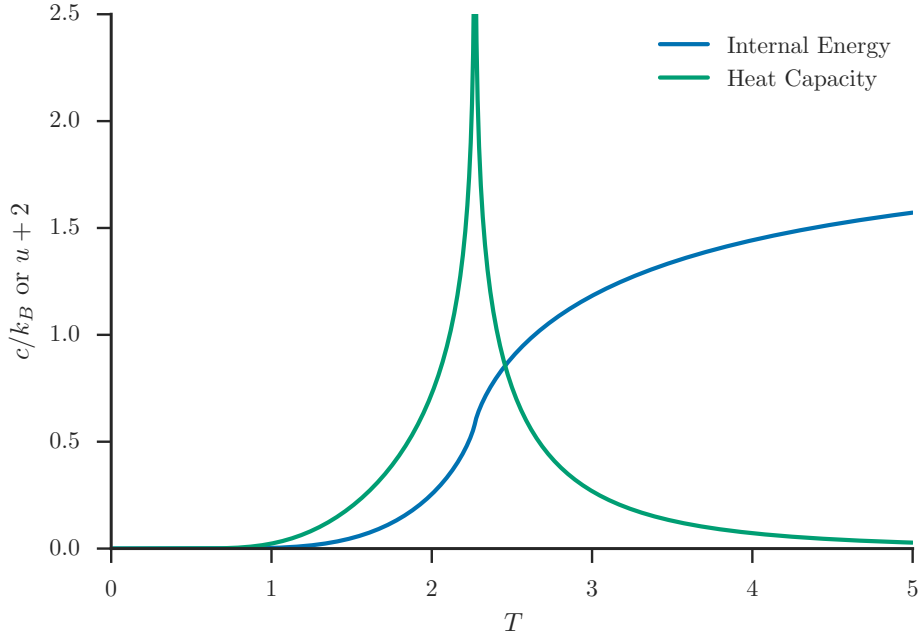


Figure 1.2: The internal energy per site and the specific heat per site of the two-dimensional Ising model in zero-field. Notice the divergence of the specific heat around $T = T_c \approx 2.269$, which is cut off in this graph due to the limited resolution.

1.2.2 The Ising Model around T_c

Given that we would like to know how the Ising model behaves at T_c , we need to expand the expressions for the internal energy and the heat capacity around this temperature. When $T \approx T_c$ the argument k of the elliptic integrals becomes approximately 1. More precisely

$$k \approx 1 - 4\beta_c^2 J^2 \left(\frac{T}{T_c} - 1\right)^2, \quad (1.48)$$

$$k' \approx 2\sqrt{2}\beta_c J \left(\frac{T}{T_c} - 1\right) \quad (1.49)$$

with $\beta_c = \frac{1}{k_B T_c}$. It is obvious that for $k = 1$ $E(k) = 1$ whereas $K(1)$ diverges since the integral becomes

$$\int_0^{\frac{\pi}{2}} \frac{1}{\cos \theta} d\theta \rightarrow \infty. \quad (1.50)$$

Nevertheless an approximation around $k = 1$ yields $K(k) \approx \log\left(\frac{4}{k'}\right)$.

At T_c the internal energy per spin u does not diverge and has the value

$$u(T_c) = \frac{-J}{\tanh(2\beta_c J)} = -\sqrt{2}J. \quad (1.51)$$

The heat capacity per spin c does diverge. Using the previous expansions

$$\frac{c(T)}{k_B} \approx \frac{8}{\pi} (\beta_c J)^2 \left[\log\left(\frac{4}{k'}\right) \right] \quad (1.52)$$

$$= \frac{2}{\pi} \log(1 + \sqrt{2})^2 \left[-\log\left|\frac{T}{T_c} - 1\right| - 1 - \frac{\pi}{4} - \log\left(\frac{\sqrt{2}}{4} \log(1 + \sqrt{2})\right) \right] \quad (1.53)$$

and we see c diverging logarithmically as $T \rightarrow T_c$. This means that the critical exponents $\alpha = 0$ for the two-dimensional Ising model. When Onsager first solved the model, it also highlighted the first instance of universality. Onsager considered the general case where the interaction energy in the x - and y -directions differed, but the logarithmic divergence depended only on the critical temperature, and not on the ratio J_1/J_2 [3]. Note that the sharp phase transition as described here only occurs when the lattice is of infinite extent. When we do not operate in the thermodynamic limit the partition function of the Ising model is analytic for all T and the specific heat has a maximum but no divergence at T_c [5].

1.2.3 Magnetisation of the Ising Model

Onsager also calculated the magnetisation of the Ising model. On the square lattice with $T < T_c$ and $J_1, J_2 > 0$ the magnetisation squared is equal to the spin-spin correlation function

$$M^2 = \lim_{N \rightarrow \infty} \langle \sigma_{00} \sigma_{0,N} \rangle = \lim_{N \rightarrow \infty} \langle \sigma_{00} \sigma_{N,N} \rangle. \quad (1.54)$$

Determining the correlation function is non-trivial, but the result is

$$M^2 = \lim_{N \rightarrow \infty} \langle \sigma_{00} \sigma_{0,N} \rangle = \left[\frac{(1 - \alpha_1^2)(1 - \alpha_2^2)}{(1 - \alpha_1 \alpha_2)^2} \right]^{1/4} \quad (1.55)$$

with

$$\alpha_1 = \frac{z_1(1 - |z_2|)}{1 + |z_2|}, \quad \alpha_2 = \frac{1}{z_1} \frac{1 - |z_2|}{1 + |z_2|}, \quad (1.56)$$

where we use z_1 and z_2 as defined in equation (1.23). The magnetisation is then (see figure 1.1 for a plot)

$$M = \left[1 - \{\sinh(2\beta J_1) \sinh(2\beta J_2)\}^{-2} \right]^{1/4}. \quad (1.57)$$

When we approach the critical temperature from below we can expand around T_c and get

$$M \propto \left[4\beta_c \left\{ \frac{J_1}{\tanh(2\beta_c J_1)} + \frac{J_2}{\tanh(2\beta_c J_2)} \right\} \left\{ 1 - \frac{T}{T_c} \right\} \right]^{1/8}. \quad (1.58)$$

Comparing with equation (1.10) we see that $\beta = 1/8$ [8].

Finally the magnetic susceptibility can be shown to diverge as $\chi \propto |T - T_c|^{-7/4}$, giving $\gamma = 7/4$. With the earlier result $\alpha = 0$, we can use the scaling laws (section 1.1.2) to determine that $\delta = 15$, $\eta = 1/4$ and $\nu = 1$ [2].

1.3 The Potts Model

The Potts model was first defined by Renfrey Potts in 1953 as a generalisation of the Ising model on the suggestion of Cyril Domb, his supervisor [9]. Domb suggested that to generalise, we can consider the spins of the model to be confined to a plane, with each spin pointing in one of q equally spaced directions, separated by an angle $\theta_n = 2\pi n/q$, with $n = 0, 1, \dots, q-1$. The nearest neighbour interaction depends only on the angle between those neighbours. The Hamiltonian for this system is

$$H = - \sum_{\langle i,j \rangle} J(\theta_{ij}), \quad (1.59)$$

with $J(\theta)$ a 2π -periodic function and $\theta_{ij} = \theta_i - \theta_j$ the angle between two spins. Domb suggested to use $J(\theta_{ij}) = -\epsilon_1 \cos(\theta_{ij})$. This is now known as the planar Potts model.[8]

Making use of a duality relation which showed that the partition function at a low temperature T was equal to that at a high temperature T^* , ($Z(T) = Z(T^*)$) Potts found the critical temperature for $q = 2, 3, 4$, by assuming that there is exactly one temperature for which $T = T^*$. This method was first used by Kramers and Wannier in 1941 [10] to determine the critical temperature of the two-dimensional Ising model, three years before Onsager's exact solution.

Potts also gave a formula for the critical temperature of a system with a different interaction strength: $J(\theta_{ij}) = -\epsilon_2 \delta(\sigma_i, \sigma_j)$, where δ is the Kronecker delta. For this system he established that for all q the transition occurs at [9]

$$\frac{x_0}{x_1} = 1 + \sqrt{q}, \quad (1.60)$$

with

$$x_0 = e^{-J_0/k_B T} \quad \text{for spins in like orientations,} \quad (1.61)$$

$$x_1 = e^{-J_1/k_B T} \quad \text{for spins in unlike orientations.} \quad (1.62)$$

For $q = 3$ we find that the critical temperature is $T_c = \frac{1}{\log(1+\sqrt{3})} \approx 0.995$ [11]. We call this system the standard Potts model. Throughout the rest of this thesis we will mainly consider this model and will simply refer to it as the Potts model. The planar and standard Potts models are related for $q = 2$ by $\epsilon_2 = 2\epsilon_1$ and for $q = 3$ by $\epsilon_2 = \frac{3}{2}\epsilon_1$. For the Potts model in two dimensions the transition at the critical point is first-order for $q > 4$ and second-order for $q \leq 4$ [8, 12]. Potts found that an earlier result by Ashkin and Teller for the discontinuities of the energy and specific heat [13], ΔE and ΔC , could be generalised to show the relationship between those quantities:

$$\sqrt{q}T\Delta C = \log\left(\frac{x_0}{x_1}\right)\Delta E. \quad (1.63)$$

Therefore transitions with a continuous energy exclude the possibility of a discontinuous heat capacity. This does however not mean that the specific heat needs to be finite at T_c .

At the critical temperature the internal energy is [3, 14]

$$u_{\text{average}} = - \left(1 + \frac{1}{\sqrt{q}}\right) J, \quad (1.64)$$

with

$$u_{\text{average}} = \frac{1}{2}(u_c^+ + u_c^-), \quad (1.65)$$

where $u_c^+ = \lim_{T \rightarrow T_c^+} \langle u \rangle$ and for a second order transition $u_c^+ = u_c^-$. For $q = 3$ (a second order transition) $u_{\text{average}} = -(1 + \frac{1}{\sqrt{3}})J \approx -1.577J$.

To properly define when the phase transition occurs we define the order parameter, which here is the magnetisation, as

$$m = \frac{1}{N} \left| \sum_k^N e^{i2\pi n_k/q} \right|, \quad (1.66)$$

with the sum running over every site of the lattice, N being the number of sites and n_k being the state of the spin at site k . This is based on the definition of the q -state Potts model as consisting of equally spaced vectors. It has the desired properties of being 1 when all spins on the lattice point in the same direction and being 0 when the spins are equally distributed over the possible states.

The Potts model has not been solved exactly, thus we do not know the exact values of the critical exponents. There are however other systems suspected to be in the same universality class as the Potts model. For $q = 2$ the model reduces to the Ising model and associated exponents (section 1.2.3). For $q = 3$ the model is conjectured to be equal to adsorbed two-dimensional adsorbed monolayers of gases on crystals, a system that can be probed experimentally and gives $\alpha = 0.36$ [14]. It is also suspected that the hard hexagon model (a two-dimensional lattice model describing a gas of non-overlapping spheres, placed on a triangular lattice so no two molecules are adjacent) is in this universality class. This model is exactly solved and has the critical exponents [3, 8]

$$\alpha = \frac{1}{3}, \quad \beta = \frac{1}{9}, \quad \delta = 14, \quad \nu = \frac{5}{6}, \quad \gamma = \frac{13}{9}, \quad \eta = \frac{4}{15}. \quad (1.67)$$

While no exact solution exists, Den Nijs has put forth conjectures for the values of the exponents [15]. We have [8]

$$2 - \alpha = \frac{2}{y_t}, \quad 1 + \frac{1}{\delta} = \frac{2}{y_h}, \quad (1.68)$$

with the thermal and magnetic exponents:

$$y_t = \frac{3(1-u)}{2-u}, \quad y_h = \frac{(3-u)(5-u)}{4(2-u)}. \quad (1.69)$$

For $q \leq 4$ [8]

$$0 \leq u = \frac{2}{\pi} \arccos\left(\frac{\sqrt{q}}{2}\right) \leq 1. \quad (1.70)$$

The critical exponents then become [3, 8]:

$$\begin{aligned} \alpha &= \frac{2(1-2u)}{3(1-u)}, & \delta &= \frac{(3-u)(5-u)}{1-u}, \\ \beta &= \frac{1}{12}(1+u), & \nu &= \frac{2-u}{3(1-u)}, \\ \gamma &= \frac{u^2-4u+7}{6(1+u)}, & \eta &= \frac{1-u^2}{2(2-u)}. \end{aligned} \quad (1.71)$$

For $q = 3$, $u = 1/3$, and these equations reproduce the exponents of equation (1.67).

Chapter 2

Simulating Lattice Models

2.1 Markov Processes and Monte Carlo Methods

While some success is to be had by using analytical methods to study lattice models and their critical exponents, often the partition function cannot be determined exactly. At that point we can use computer simulation of the models in question to study phase transitions, while having exactly solved models as a benchmark for these simulations.

We are almost always interested in the value of a thermal average of a quantity:

$$\langle X \rangle = \sum_{\alpha} X_{\alpha} p_{\alpha}, \quad (2.1)$$

with $\langle X \rangle$ a thermal average, the sum running over all configurations, each labeled by α , and p_{α} the Boltzmann probability of state α in equilibrium [2]:

$$p_{\alpha} = \frac{e^{-\beta E_{\alpha}}}{\sum_{\alpha} e^{-\beta E_{\alpha}}}. \quad (2.2)$$

While for small lattices (such as 3 by 3) the number of states is tractable (this number scales as 2^N for the Ising model and q^N for the Potts model), for larger lattices the sum can no longer be performed exactly and some clever way has to be found to sample only important states.

Instead, we randomly choose states distributed according to a probability distribution p_{α} . If we choose M random states the estimator for the quantity X is

$$X_M = \frac{\sum_{i=1}^M X_{\alpha_i} \frac{1}{p_{\alpha_i}} e^{-\beta E_{\alpha_i}}}{\sum_{j=1}^M \frac{1}{p_{\alpha_j}} e^{-\beta E_{\alpha_j}}}, \quad (2.3)$$

which becomes $\langle X \rangle$ when $M \rightarrow \infty$. We could set $p_{\alpha_i} = 1$ and choose all states with equal probability. However, often only a few states dominate the sums in equation (2.3). In a large system we will then sample a huge amount of states that contribute almost nothing to the sums [7].

To resolve this problem we can use the fact that a physical system samples the states available to it according to the Boltzmann probability. We therefore set p_{α_i} equal to equation (2.2), after which the estimator becomes

$$X_M = \frac{1}{M} \sum_{i=1}^M X_{\alpha_i}. \quad (2.4)$$

This way of sampling is called importance sampling. The states that contribute the most are also sampled most often [7].

If the M observations of X_{α_i} are statistically independent and distributed according to a Gaussian, the error on the estimate is

$$\Delta X_M = \frac{\sigma}{\sqrt{M}}, \quad (2.5)$$

with

$$\sigma = \sqrt{\text{Var}(X)} = \sqrt{\langle X^2 \rangle - \langle X \rangle^2} \quad (2.6)$$

the standard deviation. We see that the error on an estimate goes like $M^{-1/2}$, meaning that with more samples the error becomes smaller. The above equation is only valid for statistically independent measurements, which the measurements done in the simulations generally will not be [4]. Dealing with correlated measurements is discussed in section 3.1.

To pick states randomly according to the Boltzmann distribution we use a Markov process. In a Markov process a system in state μ is mutated into a state ν with probability $P(\mu \rightarrow \nu)$. These probabilities do not vary over time and depend only on the system as it is in state μ , not as it was earlier. The probabilities should satisfy

$$\sum_{\nu} P(\mu \rightarrow \nu) = 1 \quad (2.7)$$

since the system is always in some state [4, 7]. To simulate a lattice over a longer time, we look at a series of Markov processes, called a Markov Chain.

To sample the Boltzmann distribution and allow for the use of importance sampling any algorithms we use to simulate lattice systems have to conform to two conditions. First the process has to be ergodic. This means that given a starting state, any other possible state of the system should be reachable in a finite number of steps. Second, the algorithm has to display detailed balance:

$$p_{\mu} P(\mu \rightarrow \nu) = p_{\nu} P(\nu \rightarrow \mu). \quad (2.8)$$

Under these conditions the system will always tend to the desired probability distribution after a long enough time.

The probability P can be decomposed into two parts:

$$P(\mu \rightarrow \nu) = g(\mu \rightarrow \nu) A(\mu \rightarrow \nu), \quad (2.9)$$

with g the selection probability, the probability of selecting the state ν given a starting state μ and A the acceptance ratio, how likely it is to accept a given new state. To efficiently sample states in a simulation the acceptance probability should be as large as possible.

To simulate the systems using Markov chains we use Monte Carlo simulations.¹ While specific algorithms are described below, the steps of a simulation are broadly the same. First, to be able to change spins we need to prepare the system in a given state at the start of the simulation. Common choices are to prepare the system in a $T = \infty$ state where all spins are randomly aligned, or $T = 0$ where all spins are pointing in one

¹Monte Carlo simulations have their origin in the Manhattan Project and were used after World War 2 to simulate neutron transport in hydrogen bombs [7].

randomly chosen direction (in the ferromagnetic case $J > 0$, where there are multiple ground states) [7].²

Given the finite size of the simulated lattice a boundary occurs in the system. We can deal with this by applying periodic boundary conditions, wrapping the lattice around in x - and y -directions [4].

Because the probability distribution of states tends to the Boltzmann distribution only after enough time has elapsed (equation (2.8)), the system has to be given enough time to attain this state before measurements can be taken. A way to determine when this is the case is to plot the value of quantities such as the internal energy and absolute magnetisation and determine when these values stabilise.

After every step of an algorithm, measurements are performed. We measure the energy and magnetisation, as well as higher order quantities like the energy squared (necessary to calculate values of quantities such as the heat capacity). The magnetisation for the Ising model is measured by calculating

$$M = \sum_i \sigma_i, \quad (2.10)$$

with the sum running over all spins on the lattice. Because in finite systems the spin symmetry is not broken, the system magnetisation below T_c oscillates between the possible extreme values of the magnetisation when $T < T_c$ (two possible values for the Ising model, see figure 2.1). The mean of a long enough series of measurements of the magnetisation will therefore always be approximately 0. It is then more useful to measure the absolute magnetisation per spin $m = |M|/N$.

These measurements are not statistically uncorrelated, a requirement for many statistical methods. We define the autocorrelation function [2]

$$C(k) = \frac{\langle X_{\alpha_j} X_{\alpha_j+k} \rangle - \langle X_{\alpha_j} \rangle^2}{\langle X_{\alpha_j}^2 \rangle - \langle X_{\alpha_j} \rangle^2} \quad (2.11)$$

with X_{α_j} the values of X when the system is in state α at time j . The autocorrelation function is 1 when $k = 0$, meaning complete correlation between states, and is approximately 0 when k is large enough that the states are no longer correlated. For long enough time scales the autocorrelation falls off as

$$C(k) \propto e^{-k/\tau}, \quad (2.12)$$

where τ is the typical time for this decay and is called the correlation time. Because after a single correlation time the correlation function is still $1/e \approx 0.367$, it is best to sample once every 2τ steps, when the correlation is only $1/e^2 \approx 0.135$ [7].

2.2 The Metropolis Algorithm

The Metropolis algorithm, first published by Metropolis *et al.* in 1953 [16], is a simple algorithm that was first used by its creators to study the dynamics of continuously displaceable hard spheres. A step of the algorithm is as follows [2]:

1. Choose a random site on the lattice.

²In the anti-ferromagnetic case, $J < 0$, each spin wants to have a different orientation compared to its neighbours. This results in a checkerboard pattern.

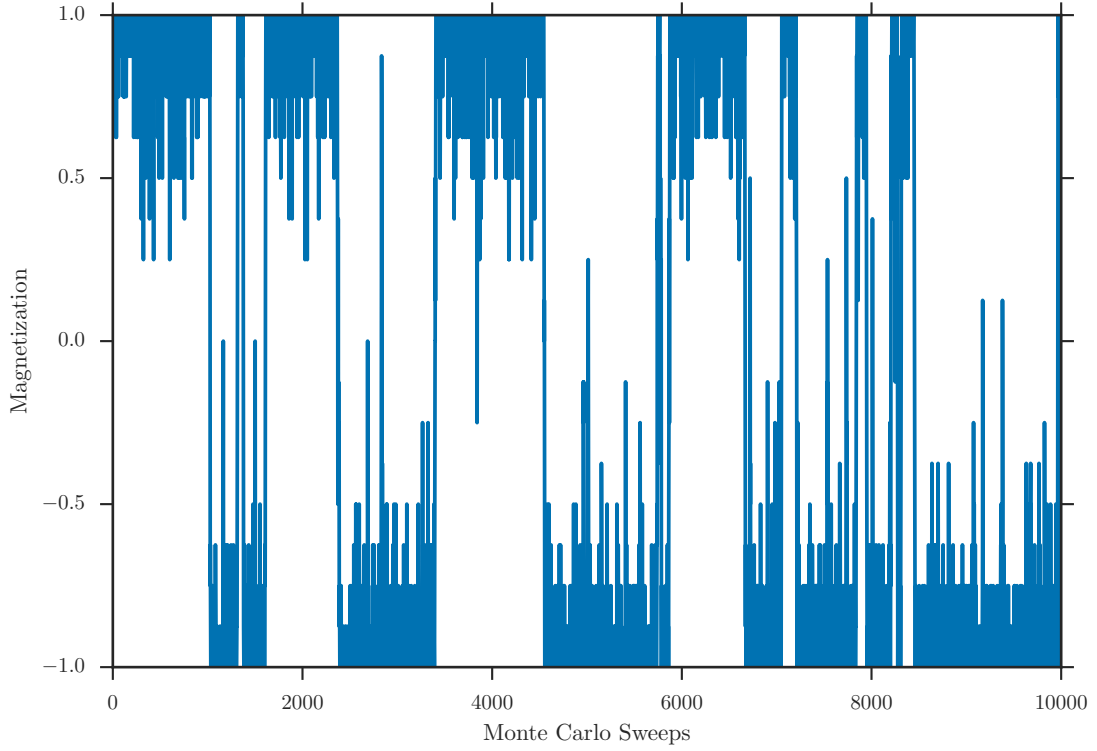


Figure 2.1: The magnetisation per spin of the Ising model. As the simulation runs (this simulation was done at $T = 2$ for a 4 by 4 system) the magnetisation oscillates.

2. Flip the spin on this location and calculate the change in energy ΔE associated with this spin-flip.
3. Calculate the probability of accepting this move:

$$A = \begin{cases} e^{-\beta \Delta E} & \text{if } \Delta E > 0 \\ 1 & \text{if } \Delta E \leq 0 \end{cases} \quad (2.13)$$

4. Generate a random variable r in the interval $[0, 1)$.
5. Accept the move if $r \leq A$. Otherwise leave the lattice as it was.

The spin flip may also happen to a non-random spin simply by iterating over the rows and columns of the lattice. When studying only equilibrium dynamics, the two approaches are identical [4]. Ergodicity is satisfied because any state can be attained by flipping a sufficient number of single spins. Detailed balance is also satisfied.

To put a measure on the time elapsed in the simulation we define a sweep. During a sweep every point on the lattice has been considered for flipping at least once. Therefore a sweep consists of N of the steps described above. The advantage of this is that time in a simulation can be measured independent of system size [7]. To keep the number of measurements reasonable, measurements are performed after every sweep, and not after every algorithm step. In figure 2.2 we see the progression towards equilibrium as a Metropolis simulation of an Ising model lattice progresses. Figure 2.3 shows how the lattice differs at equilibrium for different temperatures.

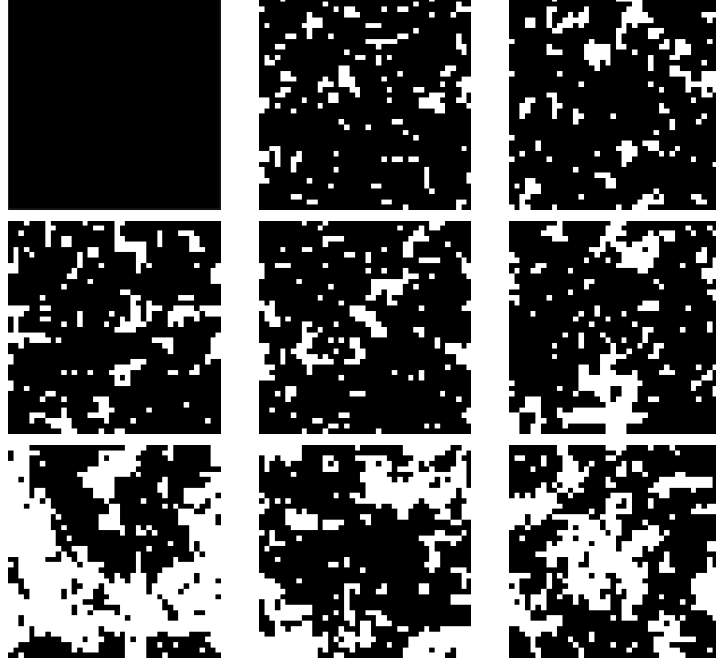


Figure 2.2: A 40 by 40 Ising model lattice simulated using the Metropolis algorithm at $T = 2.4$ after 0, 10, 20, 40, 100, 200, 400, 1000 and 5000 sweeps. The simulation started in a completely ordered $T = 0$ state.

2.2.1 Optimisations

Especially for the Ising model simulated with the Metropolis algorithm some optimisations can be included in the simulation that have a large effect on runtime. The exponentials needed for the acceptance ratio can be pre-calculated since there are (in two dimensions) only 5 possible different ways in which the neighbours are oriented: all aligned, 3 aligned, etcetera. Because the exponentials require costly floating point operations, caching those values gives a significant speed increase compared to recalculating them at every step.

For the energy of the Ising model simulated with the Metropolis algorithm there is a trick that allows for much quicker computation of the energy difference than explicitly using the Hamiltonian, which requires as many calculations as there are bonds in the system, a number that scales with N . The difference in energy between two states μ and ν is

$$E_\nu - E_\mu = -J \sum_{\langle ij \rangle} \sigma_i^\nu \sigma_j^\nu + J \sum_{\langle ij \rangle} \sigma_i^\mu \sigma_j^\mu \quad (2.14)$$

$$= -J \sum_{\substack{i \text{ nearest} \\ \text{neighbour to } k}} \sigma_i^\mu (\sigma_k^\nu - \sigma_k^\mu) \quad (2.15)$$

The term in brackets can only be -2 or $+2$, depending on the value of the spin k to be flipped before the flip. Thus $\sigma_k^\nu - \sigma_k^\mu = -2\sigma_k^\mu$ and

$$E_\nu - E_\mu = 2J\sigma_k^\mu \sum_{\substack{i \text{ nearest} \\ \text{neighbour to } k}} \sigma_i^\mu. \quad (2.16)$$

This expression depends only on the neighbours of the spin that is to be flipped and can be calculated before a spin is flipped. Using that $E_\mu = E_\nu + \Delta E$ we only have



Figure 2.3: Typical structures in a Metropolis simulation of a 20 by 20 Ising model at different temperatures after 5000 sweeps. From top left: $T = 2.2, 2.3, 2.4, 2.8, 3.4, 3.8$

to calculate the energy using the Hamiltonian once at the start, and then update the energy after every step with the change in energy. A similar trick can be used for the magnetisation, where the change in magnetisation can be shown to be $\Delta M = 2\sigma_k^\mu$ [7].

2.2.2 Critical Slowing Down

The Metropolis algorithm is very simple but does have one major drawback. Close to the phase transition, the correlation time diverges. This is because there the correlation time goes like

$$\tau \propto \xi^z, \quad (2.17)$$

where z is called the critical dynamic exponent and is different for each algorithm and system. During critical slowing down in single spin-flip algorithms the correlation length diverges (up to the system size L) and islands are formed in which all spins point in the same direction. A statistically independent configuration would need to have a significant chance of flipping the spins in such an island. However, flipping a spin whose neighbours are pointing in the same direction is energetically unfavourable and thus has a suppressed acceptance probability. Therefore the correlation time becomes large and the simulation would have to run for a long time to create enough statistically independent measurements [2], especially since at T_c we have $\xi = L$, meaning that it would quickly become impractical to simulate large systems. For the two dimensional Ising model simulated with the Metropolis algorithm the dynamic critical exponent has been measured to be $z = 2.1665(12)$ [17].

2.3 The Wolff Algorithm

The Wolff algorithm significantly improves in the area of critical slowing down, having a dynamic critical exponent of $z = 0.25(2)$ [7]. It was first proposed by Wolff in 1989 [18], and works by not flipping single spins, but building clusters of spins pointing in the same direction and flipping those en masse. The Wolff algorithm goes as follows [4, 7]:

1. Choose a random spin on the lattice i

2. Look at the neighbours of i . If a neighbouring spin j points in the same direction as i , add it to the cluster with probability $P_{\text{add}} = 1 - e^{-2\beta J}$.
3. Repeat step 2 for spin j until no more spins can be added to the cluster.
4. Flip the entire cluster.

If a spin was considered during a previous step but not added to the cluster, this spin may be reconsidered for future steps. Spins that are already part of the cluster are not added again. The algorithm fulfills the condition of detailed balance because there is always a non-zero chance that a cluster will consist of only a single spin. A succession of such single spin-flip clusters can reach any state. Detailed balance is also satisfied. Given the much smaller critical exponent for the Wolff algorithm compared to the Metropolis algorithm, critical slowing down is mostly absent.

To compare the correlation times of the Metropolis and Wolff algorithm, we need to properly define the correlation time of the Wolff algorithm. A single step will create a cluster that could correspond to a great many steps of the Metropolis algorithm. On average a Wolff cluster will cover a fraction of the lattice $\langle n \rangle / L^2$. (See figure 2.4. As the temperature increases, the probability to add a spin to the cluster decreases, meaning average cluster size decreases.) This corresponds to a single sweep of the lattice in a Metropolis simulation. The correlation time is then

$$\tau = \tau_{\text{steps}} \frac{\langle n \rangle}{L^2}. \quad (2.18)$$

This is especially important in the high temperature regime, where many single-spin clusters will form. Without the correction the correlation time would be too large [7]. Comparing correlation time for the Ising model between the Metropolis and Wolff algorithm, it is found that there is indeed a marked difference. For Wolff simulations the correlation time of the absolute magnetisation is always of order 1, even for lattices as large as 40 by 40. In contrast Metropolis simulations show a correlation time at the critical point of ~ 12 sweeps for even modest sized 10 by 10 lattices.

For the Wolff algorithm and the Ising model it is possible to calculate the change in magnetisation by considering that the change in magnetisation for a single spin is $\Delta m = -2\sigma$. The total change in magnetisation caused by flipping a spin cluster is then $\Delta m = -2\sigma \times \text{cluster size}$. This is quicker to calculate than explicitly considering every spin on the lattice to find the magnetisation.

2.3.1 Generalising Algorithms to the Potts Model

Both the Metropolis and Wolff algorithm can be easily generalised to the q -state Potts model. For the Metropolis algorithm we again pick a random spin s in state i . We then pick a new state j from the remaining $q - 1$ possible state for that spin such that $\sigma_i \neq \sigma_j$. After that we proceed as described earlier. Just like for the Ising model the algorithm satisfies the ergodicity and detailed balance conditions. For large q more efficient algorithms exist, but for $q = 3$ the Metropolis algorithm is adequate. There is still critical slowing down, with a dynamic critical exponent of $z = 2.198(8)$ [7] or $z = 2.171(62)$ [11].

Given this slowing in the Potts model cluster algorithms are again more appropriate when investigating the temperature range around the phase transition. The generalised Wolff algorithm proceeds exactly the same as in the Ising case, with the only difference

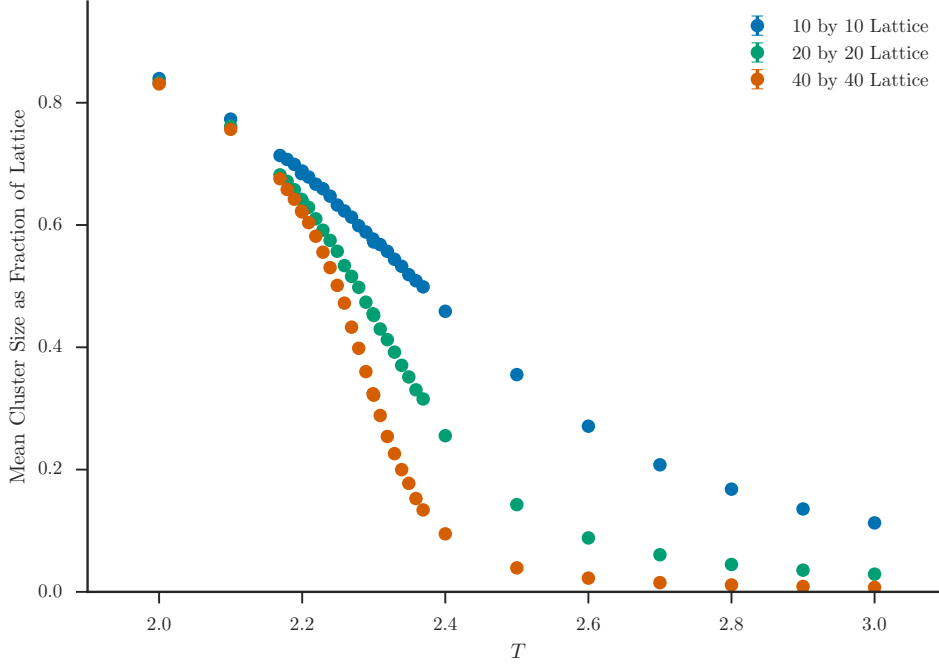


Figure 2.4: The mean size as a fraction of the lattice of clusters generated with the Wolff method. During the simulation the temperature varied from 2 to 3 in steps of 0.1, with a higher resolution of 0.01 from 2.169 to 2.369. The error bars are too small to see.

being that the probability to add a spin to the cluster has become $P_{\text{add}} = 1 - e^{-\beta J}$. We can see where this factor of 2 comes from by considering the $q = 2$ Potts model. The Hamiltonian is still

$$H = -J \sum_{\langle ij \rangle} \delta(\sigma_i, \sigma_j), \quad (2.19)$$

but for the two-state model we can rewrite this as

$$H = -\frac{1}{2}J \sum_{\langle ij \rangle} 2 \left(\delta(\sigma_i, \sigma_j) - \frac{1}{2} \right) - \sum_{\langle ij \rangle} \frac{1}{2}J. \quad (2.20)$$

These expressions are the same except for a constant, since for $\sigma_i = \sigma_j$ we have $2(\delta(\sigma_i, \sigma_j) - \frac{1}{2}) = +1$ while it is -1 otherwise. We therefore have an interaction energy $\frac{1}{2}J$ instead of J , explaining the change in the addition probability. This algorithm again satisfies detailed balance and ergodicity [7]. The dynamic critical exponent is $z = 0.60(2)$, again a significant improvement over the Metropolis algorithm.

Chapter 3

Analysing Monte Carlo Simulations

3.1 Statistics

Now that we have obtained measurements for the energy and magnetisation of the Ising and Potts models, we need to find a way to make these measurements statistically uncorrelated, since the state of a lattice after a single algorithm sweep may not be very different compared to the state before. To calculate the error on statistically independent measurements we use equation (2.5). It is possible to modify this equation for correlated measurements as

$$\Delta = \sqrt{\frac{\sigma^2}{M} \left(1 + 2\frac{\tau}{\delta t}\right)}, \quad (3.1)$$

with τ the correlation time and δt the time step between measurements [4]. In this case we need to know the correlation time of the measured quantity. It is often easier to use the Binning method, which does not require explicit knowledge of the correlation time and actually allows for the calculation of those correlation times.

3.1.1 Binning

The Binning method creates uncorrelated sequences of data from a given correlated output. Consider an original data set $A_i^{(0)}$ with $i = 1, \dots, N$ and N the number of data points in the original set. Iteratively combine two consecutive data points into a bin according to

$$A_i^{(l)} = \frac{1}{2} \left(A_{2i-1}^{(l-1)} + A_{2i}^{(l-1)} \right) \quad \text{with } i = 1, \dots, N_l = \frac{N}{2^l}, \quad (3.2)$$

with l the current binning step.¹ The average in each bin is less correlated with every step while the mean remains the same. The error on a quantity for a given binning step is

$$\Delta A^{(l)} = \sqrt{\frac{\text{Var}(A^{(l)})}{N_l}}, \quad (3.3)$$

which is the same as for uncorrelated errors (equation (2.5)). As the binning progresses the error for each step converges to the actual error, meaning that in a plot the errors

¹The original data length should be a power of two.

tend to a plateau (figure 3.1). This method can then be used to calculate the error on the energy, magnetisation and any higher order data such as E^2 .

From the binning method we can also determine the autocorrelation time [19]:

$$\tau = \frac{1}{2} \left[\left(\frac{\Delta A^{(l_{max})}}{\Delta A^{(0)}} \right)^2 - 1 \right] \quad (3.4)$$

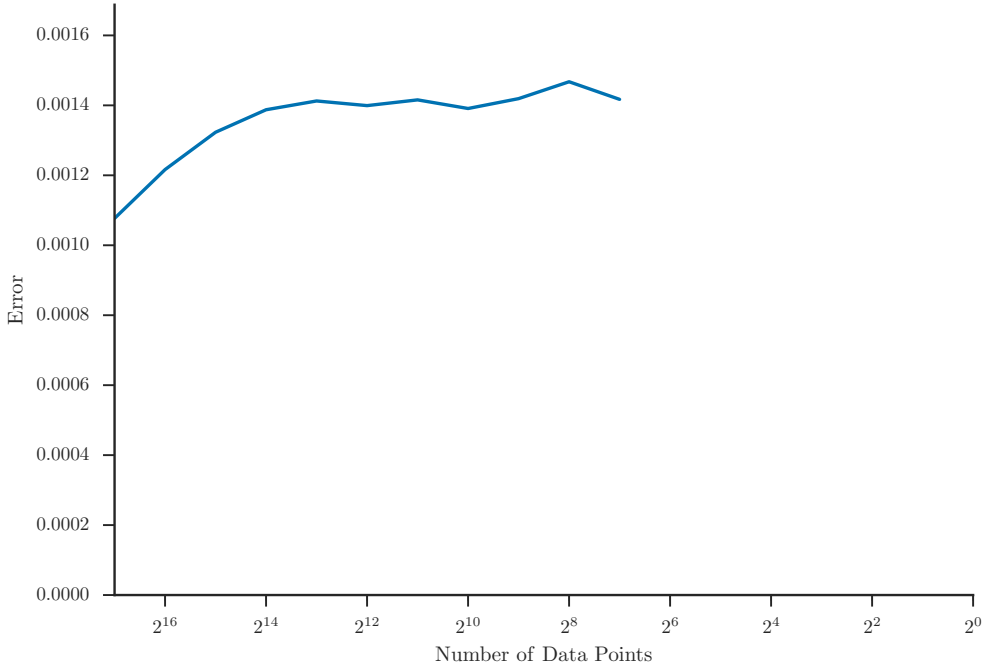


Figure 3.1: The error for the energy per site as a function of the data size. As the binning progresses the error tends to a plateau. A 4 by 4 Ising model was simulated at a temperature of 2, thermalised for 10000 sweeps and then measured for 131072 sweeps.

3.1.2 Bootstrap and Jackknife

Other interesting values we would like to know are functions of measured values. These include the heat capacity c and magnetic susceptibility χ :

$$c = \frac{\beta^2}{N} \left(\langle E^2 \rangle - \langle E \rangle^2 \right), \quad (3.5)$$

$$\chi = \beta N \left(\langle m^2 \rangle - \langle m \rangle^2 \right). \quad (3.6)$$

These quantities are not defined at a single instant of the simulation, but instead depend on time averages of the energy or magnetisation. Note that binning has to happen for both E and E^2 separately when statistically independent samples are created. To calculate the error on a quantity such as the heat capacity, we could run the same simulation a large number of times and get the error from the variance of the values

determined for c . Given the long running times of Monte Carlo simulations this is not always practical.

To still be able to calculate errors on the quantities we use resampling methods. The first method that was used is the bootstrap method. For concreteness we consider the specific heat. Given a list of n measurements of the energy (and the energy squared), we randomly choose n values, allowing duplicates. Strictly speaking these measurements should be independent, but for the bootstrap method this does not matter. We can simply use measurements made after every sweep of the lattice. With the n samples we calculate the desired quantity. Then we repeat the random sampling a number of times (1000 should be enough). With those values of the heat capacity, we can use methods for uncorrelated samples to calculate errors, such as equation (3.3) [7].

We can also use the jackknife method. Here we do need statistically uncorrelated samples. With these samples we calculate the heat capacity c_0 . Then, we calculate the heat capacity c_i for which we took out the i -th measurement from our data set. Replacing it we next remove the $i + 1$ -th measurement, and so on [7]. The error becomes [19]

$$\Delta c = \sqrt{M-1} \left(\frac{1}{M} \sum_{i=1}^M [c_i^2 - c^2] \right)^{1/2}, \quad (3.7)$$

with M the number of statistically independent measurements.

Following the advice of Landau and Binder [4], when different methods give different errors we use the largest error calculated.

3.2 Finite Size Scaling

Given these statistically independent measurements we can extract the critical exponents for different models by using finite size scaling. The advantage of this method is that T_c does not need to be known in advance.

We derive finite size scaling for the magnetic susceptibility. Close to the critical temperature we have $\chi \propto |t|^{-\gamma}$ with $t = (T - T_c)/T_c$ (equation (1.9)). We also know that for the correlation length $\xi \propto |t|^{-\nu}$ in this region (equation (1.8)). Therefore we can write $\chi \propto \xi^{\gamma/\nu}$.

The finite size of simulated systems introduced a cutoff for the correlation length equal to the system size L , also cutting off the divergence of the susceptibility. If ξ is the correlation length of the infinite system, the cutoff occurs when $\xi > L$. For $\xi \ll L$ χ should be the same as in the infinite system. We can then write for χ :

$$\chi = \xi^{\gamma/\nu} \chi_0(L/\xi), \quad (3.8)$$

where χ_0 has the properties

$$\chi_0(L/\xi) = \text{constant} \quad \text{for } L \gg \xi, \quad (3.9)$$

$$\chi_0(L/\xi) \propto (L/\xi)^{\gamma/\nu} \quad \text{as } L/\xi \rightarrow 0. \quad (3.10)$$

Because we do not know the correlation length of the infinite system it is more useful to define

$$\tilde{\chi} = (L/\xi)^{-\gamma} \chi_0((L/\xi)^\nu), \quad (3.11)$$

with which we can rewrite equation (3.8) to get

$$\tilde{\chi}(L^{1/\nu}t) = L^{-\gamma/\nu} \chi. \quad (3.12)$$

We call $\tilde{\chi}$ the scaling function. All dependence on system size has been extracted from the scaling function, meaning that its value is the same for all L when close to the critical temperature. Given a set of data for the susceptibility, we can calculate the scaling function for different γ , ν and T_c . If the values are correct, the data collapses onto a single curve around $t = 0$ when plotted (figure 3.4). For the specific heat and magnetisation similar equations can be derived [7]:

$$\tilde{c}(L^{1/\nu}t) = L^{-\alpha/\nu}c, \quad (3.13)$$

$$\tilde{m}(L^{1/\nu}t) = L^{\beta/\nu}m. \quad (3.14)$$

If T_c is known, simulations for different system sizes at T_c can be performed and a log-log plot of the data can be created to determine ratios of critical exponents. For this we use that at the critical point $t = 0$, meaning that $\tilde{\chi}$ becomes a constant independent of L , and the susceptibility scales as $\chi \propto L^{\gamma/\nu}$. Then the value of the susceptibility scales with system size as

$$\log(\chi) = \gamma/\nu \log(L) + \text{constant}. \quad (3.15)$$

Plotting the values, the slope of the line passing through the points, which can be determined by a linear fit, gives the ratio γ/ν (see figure 3.2) [19].

Two comments are in order. First, when $\alpha = 0$ the scaling function goes as $\log(L)$ [4]. This is more generally the case when any critical exponent is zero. Second, and specific to the heat capacity, we need to include a size independent background to the scaling function [20]. That is

$$\tilde{c} \propto L^{\alpha/\nu} + c_{\text{background}}. \quad (3.16)$$

Without this correction no data collapse will be possible that allows for the extraction of correct critical exponents.

3.3 The Binder Cumulant

With a finite size scaling data collapse it is possible to find the critical temperature, but fitting three parameters requires that a large parameter space is searched. An alternate way to find T_c is therefore desirable. Because of the finite size of the lattice a sharp phase transition is not present and it is not possible to determine the critical temperature by determining when the order parameter is zero. We can however define the Binder cumulant [21]

$$U_4 = 1 - \frac{\langle m^4 \rangle}{3\langle m^2 \rangle^2}, \quad (3.17)$$

which for $L \rightarrow \infty$ becomes $2/3$ for $T \rightarrow 0$ and 0 for $T \rightarrow \infty$ [4]. At the critical temperature U_4 has the same value independent of the lattice size. This is because in equation (3.14), \tilde{m} becomes a constant independent of lattice size (since $t = 0$). We

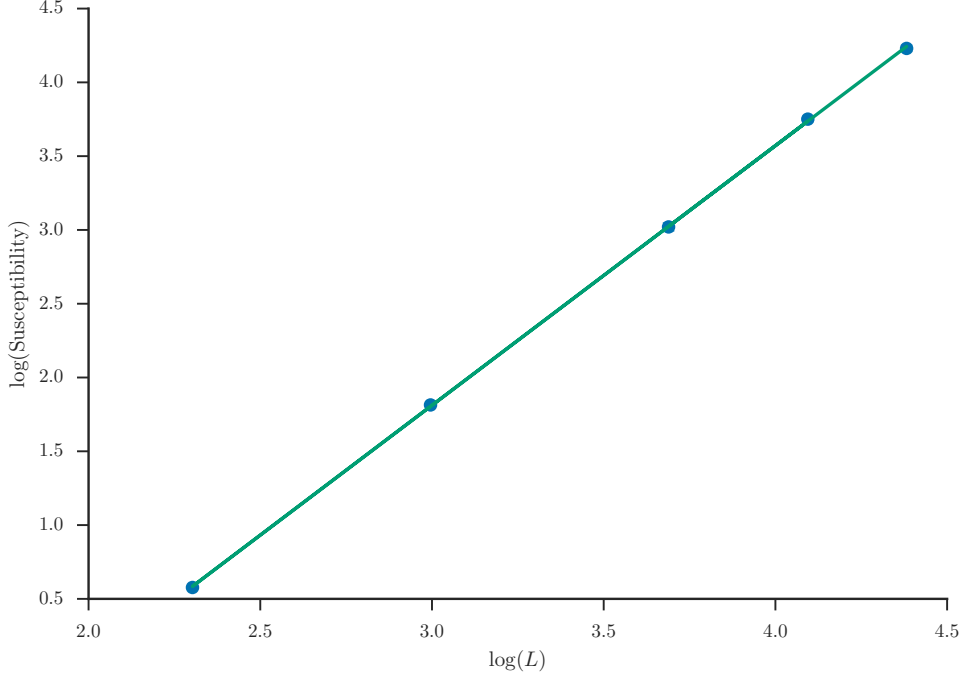


Figure 3.2: Log-log plot of the magnetic susceptibility of the Ising model at T_c for systems of size 10, 20, 40, 60 and 80. Error bars are too small to see.

have [19]:

$$U_4 = 1 - \frac{\langle m^4 \rangle}{3\langle m^2 \rangle^2} \quad (3.18)$$

$$= 1 - \frac{\tilde{m}_4(L^{1/\nu}t)L^{-4\beta/\nu}}{3\tilde{m}_2(L^{1/\nu}t)L^{-4\beta/\nu}} \Big|_{t=0} \quad (3.19)$$

$$= 1 - \frac{\tilde{m}_4(L^{1/\nu}t)}{3\tilde{m}_2(L^{1/\nu}t)} \Big|_{t=0} \quad (3.20)$$

$$(3.21)$$

which has a universal value for all lattice sizes. Therefore the critical temperature can be found by plotting the Binder cumulant for different lattice sizes and looking for the intersection (as seen in figure 3.3). If a value for the critical temperature is found in this way, it can also be plugged into the finite size scaling data collapse.

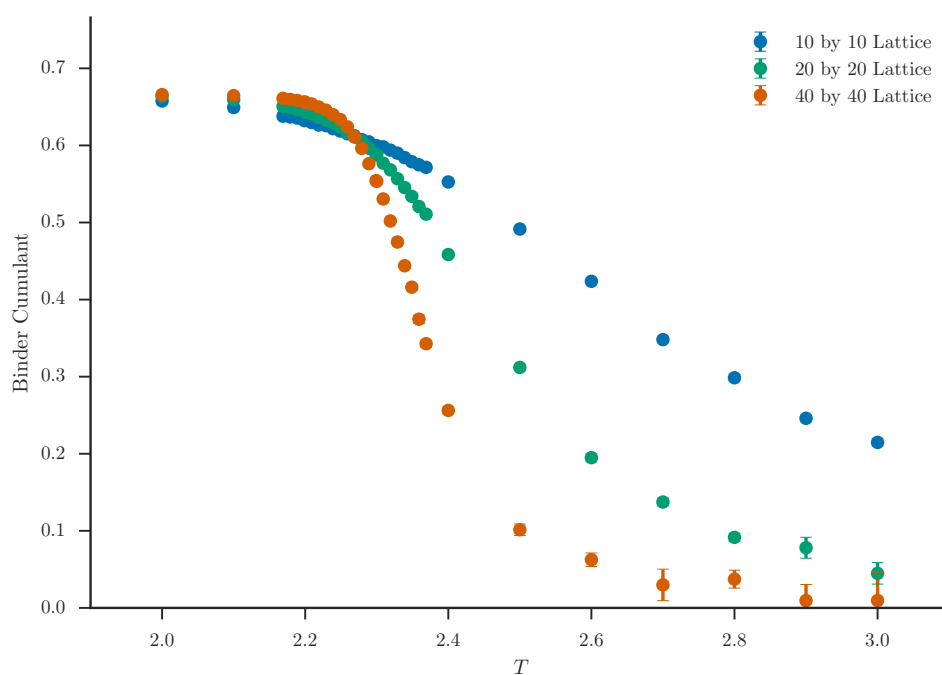


Figure 3.3: The Binder cumulant for the Ising model at three different lattice sizes. The critical point occurs at approximately $T = 2.26$. The values were acquired by simulation the lattice using the Wolff algorithm and running 65536 steps after equilibration. During the simulation the temperature was varied from 2 to 3 in steps of 0.1, with a higher resolution from 2.169 to 2.369 in steps of 0.01.

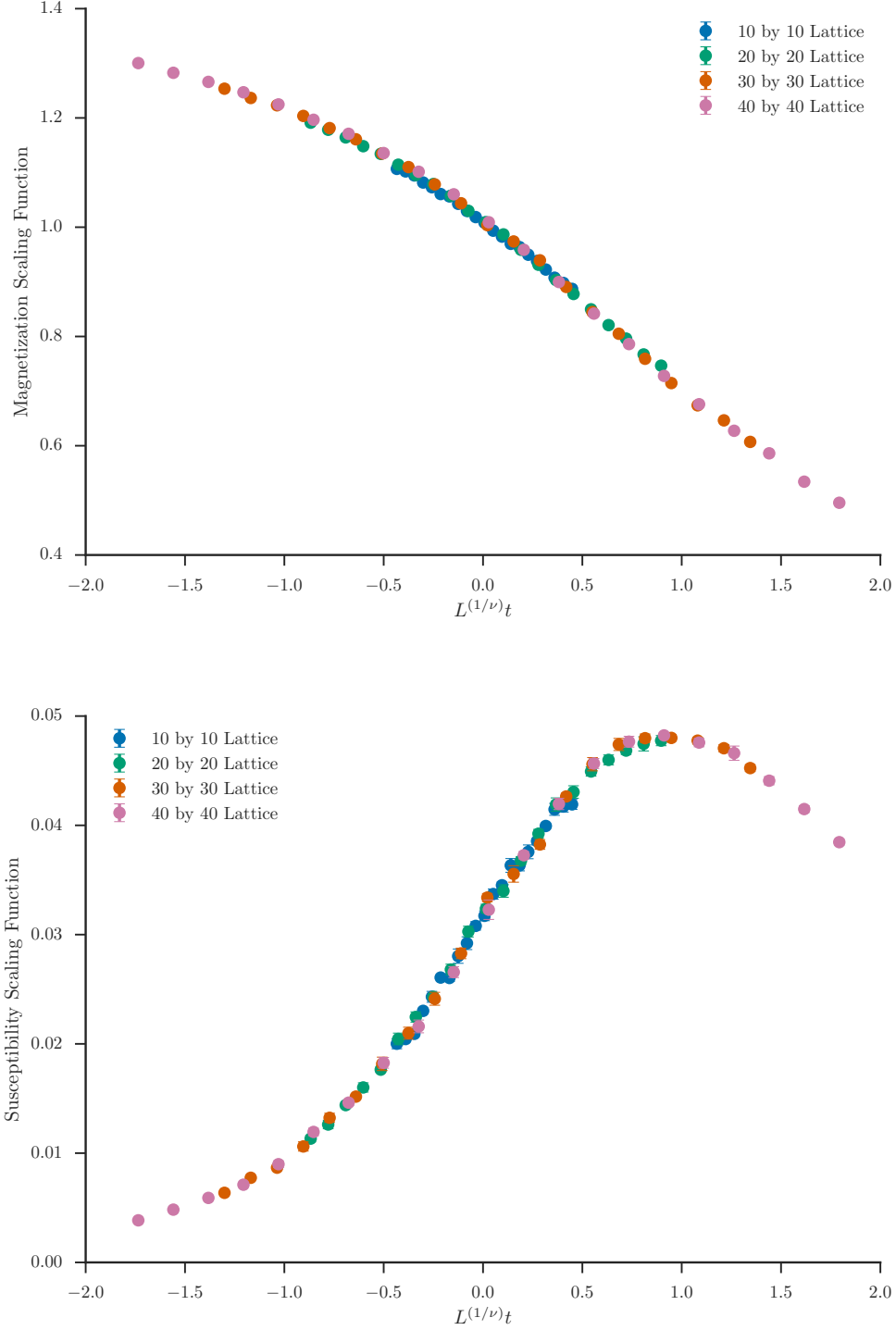


Figure 3.4: Finite size scaling data collapse for the magnetisation and the magnetic susceptibility of the Ising model. The exact values for the exponents $\gamma = 7/4$, $\nu = 1$ and $\beta = 1/8$ were used. The error bars are too small to see in these plots.

Chapter 4

Results

All results were obtained with simulations of the Wolff algorithm around the critical point. Runs consisted of 10000 equilibration sweeps, followed by 65536 measurement sweeps. For the Wolff algorithm a sweep was defined as constructing a single cluster. By binning measurements until 128 data points remained, statistically independent measurements were created.

4.1 The Ising Model at Criticality

The critical temperature of the Ising model in two dimensions in zero field was determined by finding the intersection of the Binder cumulant for lattices of sizes 10, 20, 30 and 40 (see figure 3.3, which does not include the 30 by 30 lattice). First a simulation in the temperature range 2 to 3 with steps of 0.1 was performed to roughly locate the critical point (the reported temperature all assume $k_B = 1$). A more fine-grained simulation from 2.169 to 2.369 with a resolution of 0.01 was performed afterwards. The critical temperature was determined to be 2.267(2). The critical temperature agrees well with Onsager's result of 2.269 [5], as well as those of Ghaemi *et al.* of $T_c = 2.2691(5)$ [22].

The intersection of the Binder cumulants was determined by linearly extrapolating between two consecutive data points and trying to find an intersection of those lines between consecutive lattice sizes. The average of the different intersection points was taken for the value. To determine the error on the estimate intersections were also found for the data points with their respective errors added and subtracted.

The Binder cumulant intersection for the Ising model was found to drift somewhat as different lattice sizes are considered. The intersection of lattices of size 10 by 10 and 20 by 20 and the intersection of 20 by 20 and 30 by 30 lattices was found to occur at a temperature that was lower than the critical temperature ($T_c = 2.2646(5)$ and $2.2651(3)$ respectively). The intersection between 30 by 30 and 40 by 40 lattice Binder cumulants occurred at $T_c = 2.2723(4)$, and overshoot the exact critical temperature. It would be expected that the critical temperature decreases from a value larger than the exact value and decreases as L increases. This would agree with the way the maximum of quantities like the heat capacity and magnetic susceptibility moves towards lower T as the lattice size increases.

Using log-log plots the ratio of critical exponents was determined. The ratio β/ν was determined by a linear fit of the magnetisation and was found to be 0.125(1). From a fit of the susceptibility (figure 3.2), we found $\gamma/\nu = 1.758(8)$. Both of those values agreed well with the exact values of $\beta/\nu = 0.125$ and $\gamma/\nu = 1.75$. The presented errors are as

reported by the linear regression tool used (`numpy.polyfit`).

For the heat capacity a value inconsistent with theory was found. A log-log plot gave $\alpha/\nu = 0.27(2)$, while theory predicts 0. This value was found while taking into account the background term needed in the collapse of the heat capacity. The background term was determined by finding when the finite size heat capacity data collapse was best (see below). This was the case when $c_{\text{background}} = 3$. Simply adding a background that gave a correct ratio is not a good approach, because by making the background arbitrarily large, an arbitrarily small ratio can be achieved. The data collapse would also be poor. Using the value of the constant in the expansion of the heat capacity of the Ising model around T_c from equation (1.53) ($c_{\text{background}} = 0.3$), the collapse is also very poor, with no curves intersecting.

With the determined critical temperature and ratio of critical exponents a data collapse was performed by having ν run over a range of 0.01 to 1.1 in steps of 0.01. As suggested by Sandvik [23], the quality of the data collapse was assessed using a chi least squares test. A polynomial of degree 5 was fitted to the data in the range where most data points fell after scaling, which was in the interval $L^{1/\nu}t \in [-1, 1]$. The collapse with the least spread and therefore the lowest residual per degree of freedom (the number of data points minus the number of parameters to be fitted) was then considered to produce the best fit. The values found, as well as those determined using scaling relations, are found in table 4.1. Errors were determined by adding and subtracting the error on the critical temperature and redoing the collapse, as well as adding and subtracting the errors on the ratios of critical exponents.

	α	β	γ	η	ν	δ
Exact	0	$\frac{1}{8}$	$\frac{7}{4}$	$\frac{1}{4}$	1	15
γ/ν	-0.022(8)	0.120(8)	1.762(8)	0.240(1)	1.001(4)	15.7(3.6)
β/ν	0.033(3)	0.1230(7)	1.721(4)	0.250(2)	0.983(1)	15(1.4)

Table 4.1: Critical exponents of the Ising model. The values were calculated by using the given ratio, iterating over different values of ν and determining the other critical exponent using the ratio, and then determining which values gave the best fit. The other critical exponents are calculated using scaling laws. If data points for smaller system sizes are left out, the fluctuations in the values of the critical exponents result in error bars which are potentially larger than those reported here.

For γ/ν most critical exponents agreed somewhat with the expected values, although not within their error bars, with the exception of β . The value of α is negative, which is not physical, but this stems from the value of ν being slightly too large. Given that α is still small the value seems reasonable. With β/ν the critical exponents also agreed somewhat with exact values, notably η and δ .

For γ/ν and β/ν collapses were also performed while leaving out a data set for a given data size. All data sizes were left out one by one. The drift observed in all critical exponents by doing this was always of the order 0.02 or smaller. This is larger than the values reported in table 4.1. Leaving out different data sets could thus possibly serve as an alternative way to calculate the error on critical exponents.

With the ratio α/ν it is only possible to determine α and ν , because the system of equations consisting of the scaling laws after filling in the values of α and ν is a system of three equations with four unknowns and therefore underdetermined. Using

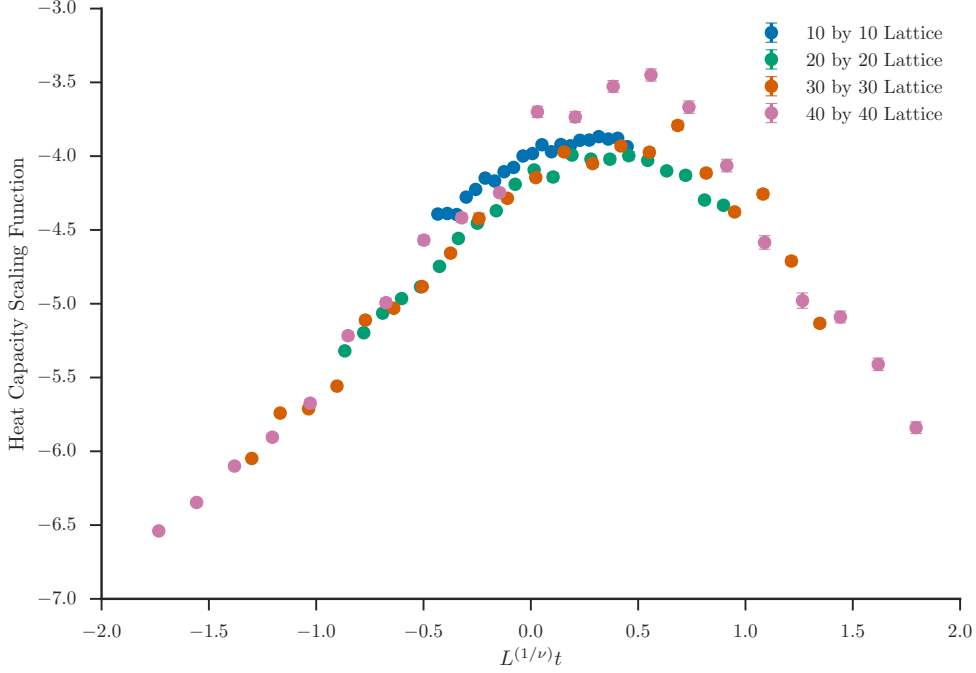


Figure 4.1: A finite size scaling data collapse for the heat capacity of the Ising model. The exact values $\alpha = 0$ and $\nu = 1$ were used. Because this corresponds to logarithmic divergence the logarithm of L was taken when calculating the scaling function. The constant background term had a value of 3.

the ratio of 0.27(2), we found $\alpha = 0.312(5)$ and $\nu = 1.096(1)$. Neither of these values agree with the exact results. The scaling laws $2\nu = 2 - \alpha$ is also not satisfied since $2.193(3) \neq 1.69(1)$. Using the exact ratio of 0 we find $\alpha = 0.003(4)$ and $\nu = 0.77(8)$. In this case the logarithm of L is used in the scaling function. The scaling laws $2\nu = 2 - \alpha$ is also not satisfied since $1.54(16) \neq 1.997(3)$. As we see in figure 4.1, the collapse for the heat capacity even with exact results is not of as high a quality as those of the susceptibility and magnetisation.

4.2 The Potts Model at Criticality

For the 3-state Potts model in two dimensions and zero-field the critical temperature was found to be $T_c = 1.020(22)$ using Binder intersection at $L = 10, 20, 40$. This is in reasonable agreement with the result of Ghaemi *et al.* [22] ($T_c = 0.9950(1)$), as well as those of Fan *et al.* [11] of $T_c = 0.9968(20)$ and the exact value of $T_c = 1/\log(1 + \sqrt{3}) \approx 0.995$. The same behaviour of the intersection temperature increasing as the system sizes increases present in the Ising model was observed here.

From a log-log plot the critical exponents ratios were found to be $\beta/\nu = 0.131(5)$, $\alpha/\nu = 0.41(2)$ and $\gamma/\nu = 1.80(3)$. The conjectured exact values for these ratios are $\beta/\nu = 2/15 \approx 0.133$, $\alpha/\nu = 2/5 = 0.4$ and $\gamma/\nu = 26/15 \approx 1.73$. The values for β/ν and α/ν are in good agreement, with γ/ν less so. Note that these values were calculated with simulations done at $T_c = 0.995$, the exact critical temperature.

	α	β	γ	η	ν	δ
Den Nijs (1979) [3, 15]	$\frac{1}{3}$	$\frac{1}{9}$	$\frac{13}{9}$	$\frac{4}{15}$	$\frac{5}{6}$	14
Blöte (1981) [24]	$\sim \frac{1}{3}$					~ 14
Fan (2007) [11]	0.396(66)	0.108(4)	1.416(62)	0.265(19)	0.816(27)	14.1(1.1)
Hu (1980) [25]	0.3071					
Straley (1973) [26]	0.05(10)	0.103(10)	1.5(1)			
Binder (1981) [14]	~ 0.4	~ 0.1	~ 1.45			
β/ν	0.34(12)	0.11(1)	1.44(14)	0.263(33)	0.83(6)	14.2(5.4)
γ/ν	0.60(6)	0.07(6)	1.26(5)	0.20(1)	0.70(3)	19.5(16.8)

Table 4.2: The critical exponents of the three-state Potts model. The values quoted for Den Nijs are conjectured exact values. For Blöte the results had to be read from a graphs, and thus do not have error margins. Hu did not report any error margins. Table VI in [8] contains more values for the critical exponents, from both numerical and analytical methods. The final two rows of values were calculated by using the given ratio, iterating over different values of ν and determining the other critical exponent using the ratio, and then determining which values gave the best fit. The other critical exponents are calculated using scaling laws. The presented values were found with a collapse done with $T_c = 0.995$.

Using the value for the critical temperature found with the Binder cumulant produced poor collapses (see figure 4.2). Therefore all other collapses were performed with the exact value of the critical temperature $T_c = 0.995$. In the collapse the error on the critical temperature was used as determined by the Binder cumulant.

From least square fits the values of the critical exponents were found (table 4.2). Here most data points fell in the interval $L^{1/\nu}t \in [-2, 2]$. With β/ν , the value for all critical exponents agreed well with the conjectured values as well as those of Fan *et al.* [11].

For the values determined with the ratio γ/ν agreement with conjectured other experimental values was poorer, with only β agreeing within error bars. For δ the error is extremely large. This is because during error propagation the ratio $\Delta\beta/\beta$ is approximately 1. Multiplying by the value of δ then gives a large error. Given the size of the error this value should not be judged as sensible.

By leaving out data sets for given system sizes, an estimate could be made how well behaved the collapse is. From table 4.3 it is clear that some critical exponents show a large drift. This table lists the critical exponents determined using the ratio γ/ν . The values drift as different data sets are left out. This drift is most apparent for α , γ and ν . Leaving out different lattice sizes may be considered as an alternate way of assessing errors on the measurements.

From the ratio α/ν it was again only possible to determine α and ν . We found $\alpha = 0.36(2)$ and $\nu = 0.89(4)$. The phenomenological constant added to the heat capacity was determined to be 1.5, which was also used in the log-log plot mentioned above. Both values agree somewhat with the conjectured and Monte Carlo values of other sources, especially the value for α . Using the final scaling law ($2\nu = 2 - \alpha$) as a consistency check it was found that these values were not consistent within error bars since $1.78(8) \neq 1.64(2)$.

For the finite size scaling collapse it is important to note that the collapse is only

Size left out	α	β	γ	η	ν	δ
None	0.60(6)	0.07(6)	1.26(5)	0.20(1)	0.70(3)	19.5(16.8)
10	0.05(6)	0.09(6)	1.76(6)	0.20(1)	0.97(3)	20(27)
20	0.23(11)	0.1(1)	1.6(1)	0.20(1)	0.88(5)	20(25)
40	0.02(7)	0.10(7)	1.78(6)	0.20(1)	0.99(4)	19(59)

Table 4.3: The critical exponents of the three-state Potts model when a data set for a given lattice size are left out, determined with the ratio γ/ν . A noticeable drift is present for the values of α , γ and ν . The values were calculated by using the given ratio, iterating over different values of ν and determining the other critical exponent using the ratio, and then determining which values gave the best fit. The other critical exponents are calculated using scaling laws. The presented values were found with a collapse done with $T_c = 0.995$.

good very close to $t = 0$. This also means that T_c has to be known to a high degree of precision (as seen above for the Potts model). Another problem is that the scaling functions are only approximately correct even when t is small. The scaling functions are in better agreement as the system size increases, but for small L correction terms to the scaling functions have to be taken into consideration [7].

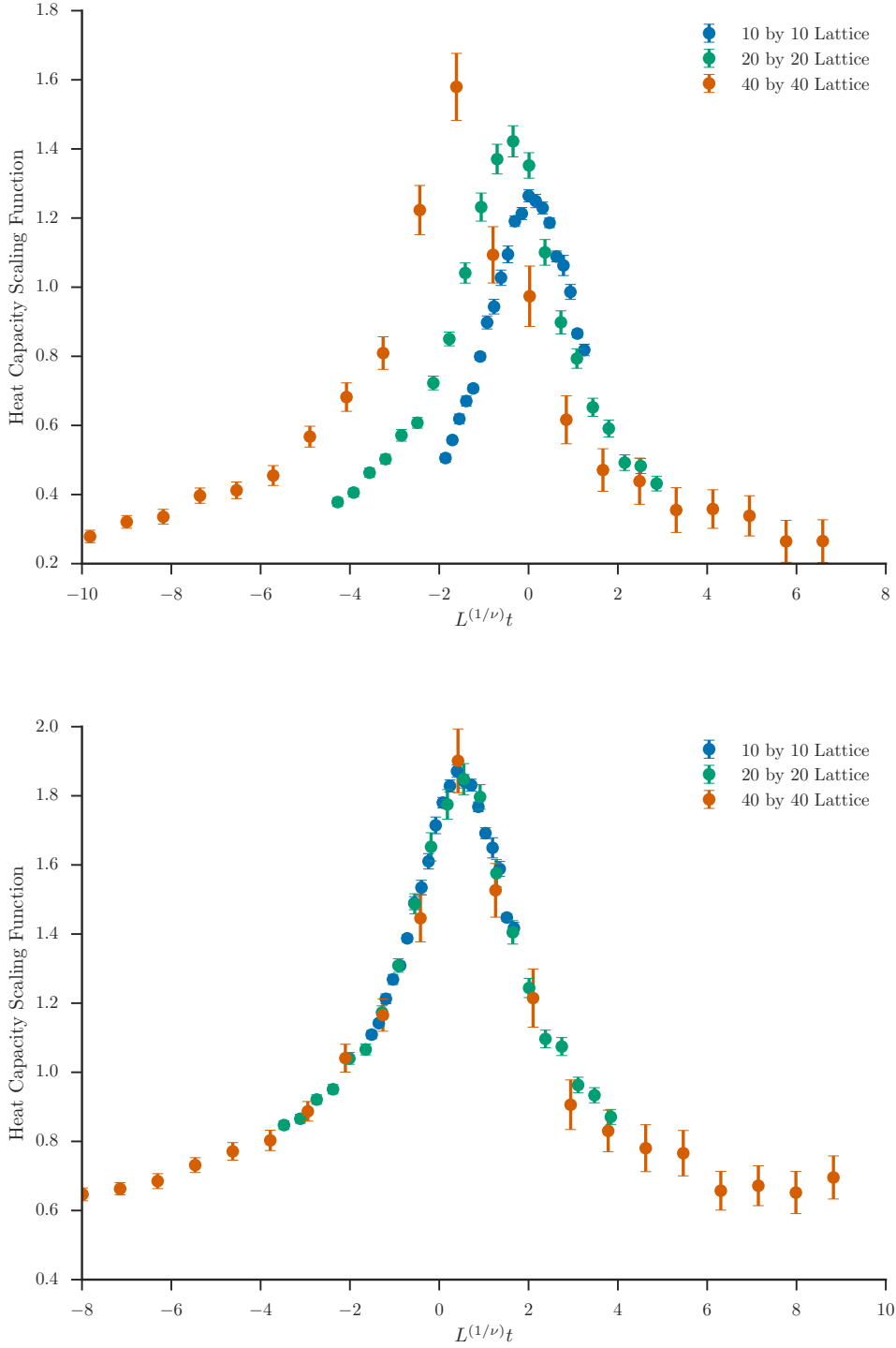


Figure 4.2: Finite size scaling collapse of the heat capacity of the Potts model. The first image uses $T_c = 1.002$, while the second one uses $T_c = 0.995$. Although the difference in critical temperature is small, the second collapse is much better. In both cases the conjectured critical exponents $\alpha = 1/3$ and $\nu = 5/6$ were used.

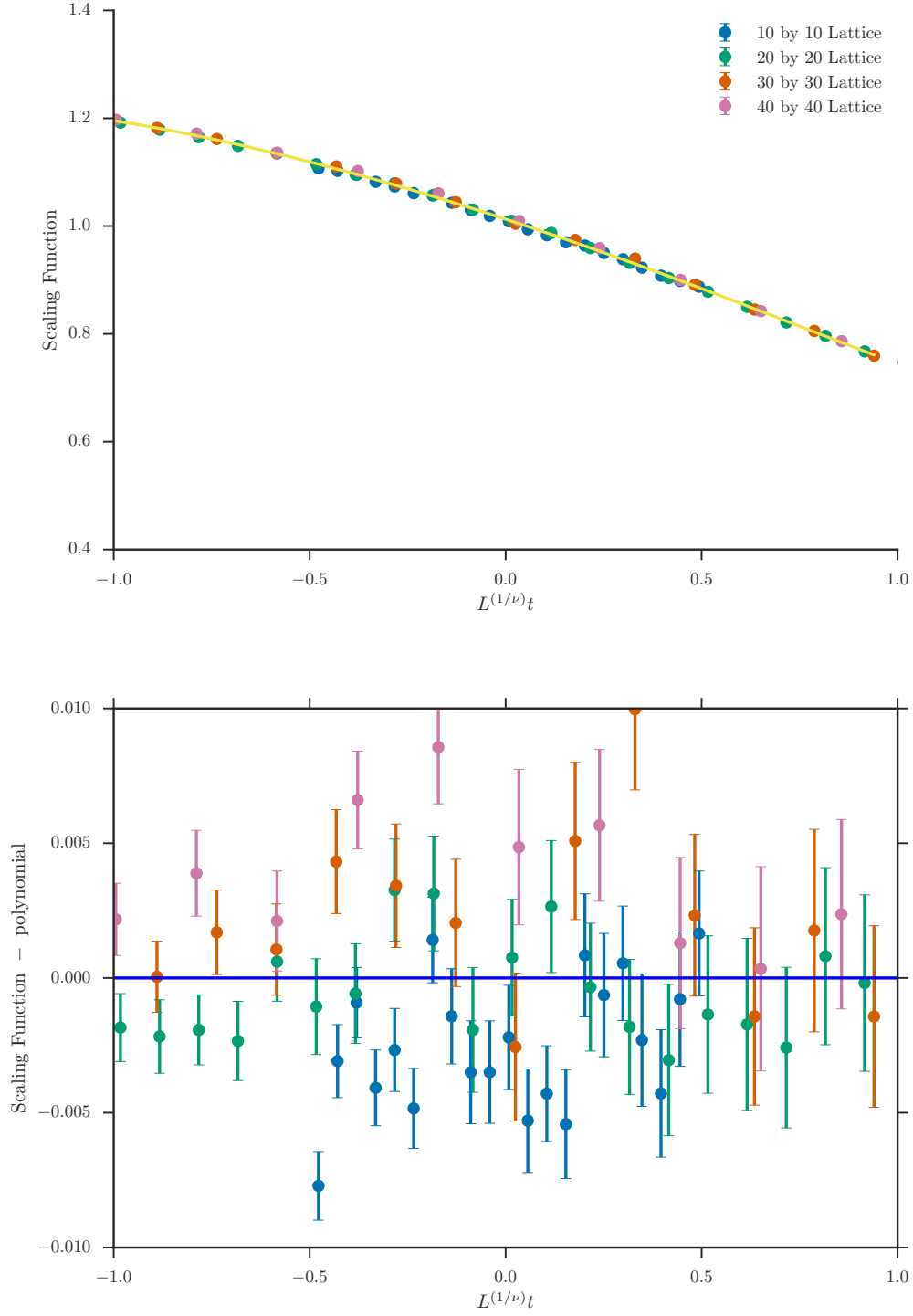


Figure 4.3: Data collapse of the magnetisation of the Ising model, using a chi least squares method to determine the goodness of fit (yellow line). The image shown represents the best collapse that could be performed. The bottom image shows the data points with the polynomial subtracted, to assess how good the fit is.

Chapter 5

Conclusions

Phase transitions and critical phenomena play an important role in life. These transitions are characterised by critical exponents, which are related to each other by scaling laws. Because only a few systems have been solved exactly, the exact values of the critical exponents of many systems are not known. In this work the critical exponents and critical temperature of the Ising model and 3-state Potts model in two dimensions and zero field were determined. To this end lattices were simulated with Monte Carlo methods, using the Metropolis algorithm as well as the Wolff algorithm. The data was analysed using finite size scaling methods. The intersection of Binder cumulants gave a value for the critical temperature, while data collapses for simulations of different sized lattices were used to determine critical exponents.

For the Ising model the value of the critical temperature as well as the ratios of critical exponents, with the exception of α/ν , agreed with the exact values of Onsager's solution. Using a data collapse the values of the critical exponents themselves were found to usually be in reasonable agreement with the exact values. The values found for α were close to zero, although one value was negative, a non-physical result. Because the error analysis with data collapses poses some problems, some of the errors reported may be underestimated. If alternatively the error bars are calculated by leaving out data sets for smaller system sizes, the error is larger and α would be 0 within error bars.

The Potts model results gave a value for the critical temperature that was in moderate agreement with the exact result. Ratios of critical exponent ratios agreed well with conjectured results, with the exception of γ/ν , which was too large. The data collapse was done using the exact value for the critical temperature as opposed to the value found using the Binder cumulant because that value gave very poor collapses. For the collapse done with β/ν all critical exponents agreed with conjectured results and the most recent Monte Carlo estimates found in the literature within error bars. The collapse using γ/ν was much poorer, with only β agreeing with conjectured results and an error on δ that was approximately as large as the value found. The values for α and ν found from α/ν were not in agreement with conjectured results.

5.1 Further Work

There are several ways to expand on the work presented here. One way is to use renormalisation group techniques to calculate the critical exponents. This is the way most papers determine those and has as an advantage that the errors on the critical exponents can be better determined.

Another approach is to study finite size scaling in more detail and try to determine the corrections that become important for small systems or far away from T_c . We could also study larger systems, for which the corrections are less important. This would require a rewrite of the simulations in a compiled language to improve execution time. It is possible that with larger system sizes the drift of quantities like the Binder cumulant could be better characterised. With larger and more system sizes, it would also be possible to find the change in critical exponents as data sets are left out while determining the critical exponents from a data collapse. This would allow for a different way of assessing errors on the critical exponents.

The Ising and Potts model can also be simulated on different kinds of lattices such as triangular or honeycomb lattices, to verify that the critical exponents (but not the critical temperature) remain the same, thus testing universality.

The autocorrelation times were determined using the data from the binning analysis. The correlation times could be investigated using an explicit calculation instead and used to establish the occurrence of critical slowing down as well as determining the dynamic critical exponents.

It is also possible to go to $q > 4$ and try to determine that the phase transition becomes first order in this case. To that end the latent heat would have to be accurately determined, while also having to deal with phase transitions that are not sharp in finite systems.

Lastly, the order parameter of the Potts model could be more closely examined. In this thesis it was defined based on the definition of the Potts model. Many papers used a slightly different definition that does not have the property of being 1 below T_c and 0 above T_c [8, 11, 14].

5.2 Acknowledgements

I would like to thank my supervisor, dr. Philippe Corboz, for introducing me into the field of computational physics and patiently explaining statistical analysis of Monte Carlo data. I would also like to thank Edan Lerner for acting as my second assessor.

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