Phase transition in compressible Ising systems at fixed volume

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Using a Ginzburg-Landau model, we study the phase transition behavior of compressible Ising systems at constant volume by varying the temperature $T$ and the applied magnetic field $h$. We show that two phases can coexist macroscopically in equilibrium within a closed region in the $T-h$ plane. It occurs near tricriticality. We find a field-induced critical point, where there is a significant change in the correlation length. The difference of the coexisting two phases and the surface tension vanish, but the isothermal magnetic susceptibility does not diverge in the mean field theory. We also investigate phase ordering numerically.

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I. INTRODUCTION

Solids are under the influence of elastic constraints and their phase transitions are often decisively influenced by couplings of the order parameter and the elastic fields. Such elastic effects strongly depend on the nature of the coupling and their understanding is crucial in technology. In the present work, we will focus on the phase transition behavior of compressible ferromagnets or antiferromagnets, which has long been studied theoretically in the physics community. In real materials, the short-range spin interactions depend on the distances among the spins, so the spin fluctuations are coupled to the elastic dilation strain. In the literature on this problem, the main issue has been the effect of the elastic coupling on the critical behavior of the spin system. A remarkable but subtle result of the renormalization group calculations is that the cubic elastic anisotropy becomes increasingly important on approaching the critical point (which is determined in the absence of the anisotropy). This renormalization effect should trigger a first order phase transition sufficiently close to the critical point. Simulations have been performed on compressible Ising systems at constant volume $V$ and a number of numerical results still remain not well understood. These theories and simulations show that the phase transition depends on whether the pressure or the volume is fixed.

In this paper, we will present a mean field theory of compressible Ising systems at constant volume using a Ginzburg-Landau free energy. Our main objectives are to demonstrate the presence of unique two phase coexistence near the tricritical point and to examine phase ordering after changing the temperature. Though our theory is a rough approximation, it will provide overall phase behavior for general values of the parameters.

The organization of this paper is as follows. In Sec. II, we will present a model, in which the order parameter and the elastic field are coupled, and eliminate the elastic degrees of freedom assuming the mechanical equilibrium condition. In Sec. III, we will examine the phase behavior in the plane of the temperature $T$ and the ordering field $h$. Detailed calculations will also be given on the susceptibility, the correlation length, and the surface tension. The presence of a unique field-induced critical point will also be reported. In Sec. IV, we will numerically integrate the time-dependent Ginzburg-Landau equation in two dimensions (2D). In the appendix, we will derive the free energy at constant pressure (or applied stress), where two-phase coexistence can be realized only on lines in the $T-h$ plane.

II. THEORETICAL BACKGROUND

A. Ginzburg-Landau free energy

We assume that a single-component order parameter $\psi$ is coupled to the elastic displacement $u$. We set up the Ginzburg-Landau free energy functional $F = F(\psi, u)$ in the form:

$$F = \int dr \left[ f_0 + \frac{C}{2} \left| \nabla \psi \right|^2 + \alpha \psi^2 \epsilon_1 + f_{el} \right],$$

(2.1)

where the space integral is within the system with volume $V$. The first part $f_0 = f_0(\psi)$ depends on $\psi$ as:

$$f_0 = \frac{\tau}{2} \psi^2 + \frac{C}{4} \psi^4 + \frac{\nu}{6} \psi^6 - h\psi.$$

(2.2)

The coefficient $\tau$ depends on the temperature $T$ as:

$$\tau = A_0 (T - T_0),$$

(2.3)

where $A_0$ is a positive constant and $T_0$ is the critical temperature in the absence of the elastic coupling. The other coefficients are treated to be independent of $T$. We fix the other field variables such as the hydrostatic pressure. The coefficients $\nu$ and $C$ are positive, while $\epsilon_1$ can be either positive or negative. The $h$ represents a magnetic or electric field conjugate to $\psi$. For antiferromagnetic materials, no uniform field conjugate to the antiferromagnetic order can be realized, so $h = 0$. We may assume $h \geq 0$ without loss of generality. If $h = 0$, $F$ is invariant with respect to $\psi \rightarrow -\psi$. The $\alpha$ represents the strength of the coupling between $\psi^2$ and the dilation strain:

$$\epsilon_1 = \nabla \cdot u.$$

(2.4)
This coupling arises when the interaction among the fluctuations of $\psi$ depends on the local lattice expansion or contraction.

In cubic crystals, the elastic energy density is of the form,

$$f_{el} = \frac{C_{11}}{2} \sum_i \epsilon_{ii}^2 + \sum_{i \neq j} \left[ \frac{C_{12}}{2} \epsilon_{ij} \epsilon_{ji} + C_{44} \epsilon_{ij}^2 \right],$$  \hspace{1cm} (2.5)

where $C_{11}$, $C_{12}$, and $C_{44}$ are the usual elastic moduli assumed to be constant, and $\epsilon_{ij} = (\nabla_i u_j + \nabla_j u_i) / 2$ is the symmetrized strain tensor. The dependence of the elastic moduli on $\psi^2$ can be important at low temperatures, however. Hereafter $(\hat{\epsilon})^2 = (\hat{\epsilon}^2)$. The elastic stress tensor $\sigma_{ij}$ is expressed as

$$\sigma_{ii} = (C_{11} - C_{12}) \epsilon_{ii} + C_{12} \epsilon_{ii} + \alpha \psi^2,$$

$$\sigma_{ij} = 2C_{44} \epsilon_{ij} \quad (i \neq j).$$  \hspace{1cm} (2.6)

Nonvanishing $\psi^2$ gives rise to a change in the diagonal stress components. We then obtain $\sum_j \nabla_j \sigma_{ij} = -\delta F/\delta u_i$, where $\psi$ is fixed in the functional derivative of $F$ with respect to $u_i$. Note that a constant hydrostatic pressure $p_0$ can be present in the reference state, where the total stress tensor is $p_0 \delta_{ij} - \sigma_{ij}$.

### B. Elimination of elastic field at fixed volume

The elastic field $u$ is determined by $\psi$ under the mechanical equilibrium condition,

$$\sum_j \nabla_j \sigma_{ij} = 0.$$  \hspace{1cm} (2.7)

Furthermore, in this paper, we impose the periodic boundary condition on $\delta u$ in the region $0 < x, y, z < V^{1/d}$. This can be justified when the solid boundary is mechanically clamped. See Appendix A for the case of fixed applied pressure. The space averages of the strains then vanish; for example, $\langle \epsilon_{ii} \rangle$. Hereafter $\langle \cdot \cdot \cdot \rangle = \int d\mathbf{r} \langle \cdot \cdot \cdot \rangle / V$. The following procedure of eliminating the elastic field has been derived by many authors in the literature in physics and engineering.\textsuperscript{1,2,4,5,11,14}

It is convenient to use the Fourier transformation, $u_j(r) = \sum_k u_j \mathbf{k} \exp(\mathbf{i}\mathbf{k} \cdot \mathbf{r})$, where $\mathbf{k}$ is the wave vector. Then the Fourier component of $\epsilon_{ii}$ is expressed as

$$\epsilon_{ii} = -\alpha \varphi_{i\mathbf{k}} / [C_{12} + C_{44} + C_{44} \zeta(\mathbf{k})],$$  \hspace{1cm} (2.8)

where $\varphi_{i\mathbf{k}}$ is the Fourier component of the variable,

$$\varphi(r) = \psi^2 - \langle \psi^2 \rangle.$$  \hspace{1cm} (2.9)

The space average of $\varphi$ is made to vanish. The $\zeta(\mathbf{k})$ is a function of the direction of the wave vector $\mathbf{k} = \mathbf{k}^{-1} \mathbf{k}$ and is defined by

$$\zeta(\mathbf{k})^{-1} = \sum_j \mathbf{k}^2_j / (1 + \xi_a \mathbf{k}^2_j),$$  \hspace{1cm} (2.10)

where $\xi_a$ is the degree of cubic anisotropy,

$$\xi_a = (C_{11} - C_{12}) / C_{44} - 2.$$  \hspace{1cm} (2.11)

We have $\zeta(\hat{\mathbf{k}}) = 1$ in the isotropic elasticity $\xi_a = 0$. After some calculations, we may eliminate $\mathbf{u}$ in $F$ to obtain the free energy $F = F(\psi)$ of $\psi$ only in the form\textsuperscript{11,13,15,16},

$$F = \int \mathbf{d}r \left[ f_0 \mathbf{C} \nabla \psi^2 - \frac{1}{2} \sum_k w(\mathbf{k}) |\varphi_{i\mathbf{k}}|^2 \right].$$  \hspace{1cm} (2.12)

The second term on the right hand side arises from the elastic coupling and is negative, where

$$w(\mathbf{k}) = \alpha^2 / [C_{12} + C_{44} + C_{44} \zeta(\mathbf{k})].$$  \hspace{1cm} (2.13)

The functional derivative of $F$ is performed to give

$$\frac{\delta F}{\delta \psi} = f_0 - C \nabla^2 \psi + 2 \alpha e_1 \psi,$$

where $f_0$ is the $F$ of $\psi$ and the Fourier transformation of $e_1$ in Eq.(2.8). In equilibrium we require $\delta F/\delta \psi = 0$.

We further simplify our free energy. In the isotropic elasticity, $w(\mathbf{k})$ is a constant independent of $\mathbf{k}$ and $e_1 = -\alpha \varphi / C_{11}$. Then $F$ is rewritten as

$$F = \int \mathbf{d}r \left[ f_0 \mathbf{C} \nabla \psi^2 - \frac{\beta}{4} (\psi^2 - \langle \psi^2 \rangle)^2 \right],$$  \hspace{1cm} (2.15)

where $\varphi$ is explicitly written in terms of $\psi$ and $\beta$ is a positive constant defined by

$$\beta = 2 \alpha^2 / C_{11}.$$  \hspace{1cm} (2.16)

The presence of the space average $\langle \psi^2 \rangle$ is a unique aspect arising from elasticity.

In cubic solids with $\xi_a < 0$, $w(\mathbf{k})$ is maximized along one of the principal crystal axes (say, along the [100] direction in 3D). If $\xi_a > 0$, it is maximized for $k_2^2 = 1/d$ for all $j$ (say, along [111] in 3D). Let $w_M$ be the maximum of $w(\mathbf{k})$ attained along these soft directions; then,

$$w_M = \alpha^2 / C_{11} \quad (\xi_a < 0),$$

$$= \alpha^2 / [K + (2 - 2/d) C_{44}] \quad (\xi_a > 0).$$  \hspace{1cm} (2.17)

where $K = C_{11} / d + C_{12} (1 - 1/d)$ is the bulk modulus. In 2D, $w(\theta) = w(\mathbf{k})$ is a periodic function of the angle $\theta$ defined by $k_x / k = \cos \theta$ and $k_y / k = \sin \theta$ with period $\pi / 2$, as displayed in Fig. 1. In phase ordering processes, the interface normals tend to be parallel to these soft directions, resulting in cuboidal domains.\textsuperscript{1,13,15,16} If the spatial inhomogeneity is mostly along these soft directions except for the edge regions of the domains, the free energy is approximately given by Eq.(2.15) with

$$\beta = 2w_M.$$  \hspace{1cm} (2.18)
existence of a single phase only. Note that $C\kappa$ of $\beta\psi$ region. To calculate it, we superimpose plane wave fluctuations of the Fourier component $\psi_k$ in the bulk region. To calculate it, we superimpose plane wave fluctuations of $\psi$ on the homogeneous average. The increase of the free energy in the second order yields $S_k$ in the Ornstein-Zernike form

$$S_k = 1/C(k^2 + \kappa^2),$$

where $\kappa$ is the inverse correlation determined by

$$C\kappa^2 = \partial^2 f_0/\partial \psi^2 - 3/\beta\psi^2 + 3\beta/\psi^2 = \tau + (3\bar{u} - 2\beta)\psi^2 + 5v\psi^4.$$  
(2.22)

In the second line, we have set $\langle \psi^2 \rangle = \psi^2$ because of the existence of a single phase only. Note that $C\kappa^2$ in the second line of Eq.(2.22) is smaller than $\chi^{-1}$ in Eq.(2.20) by $2/\beta\psi^2$. In cubic solids, $\kappa$ represents the inverse correlation length for the fluctuations varying in the softest directions. Let $\tau$ take a small negative value at $h = 0$ in the case $\bar{u} > 0$; then, $\psi^2 \approx |\tau|/\bar{u}$ from Eq.(2.19), leading to $C\kappa^2 \approx 2(1 - \beta/\bar{u})|\tau|$ from Eq.(2.22). The positivity of $\kappa^2$ is attained only for $\beta < \bar{u}$. Obviously, the disordered phase with $\psi = 0$ is unstable for $\tau < 0$. The ordered phase with $\psi^2 = -\bar{u}/2v + \sqrt{\bar{u}^2/4v^2 - \tau}$ (which is the solution of Eq.(2.19) at $h = 0$) becomes unstable for $\tau > \tau_{in}$. In particular, as $h \to 0$, we find

$$\lim_{h \to 0} \tau_{in} = -(\beta^2 - \bar{u}^2)/4v.$$  
(2.23)

III. TWO PHASE COEXISTENCE

A. Two phase states

We show that two phases can coexist in a temperature window $\tau_c - \tau_w < \tau < \tau_c$ if the parameter,

$$u = \bar{u} - \beta,$$  
(3.1)

is negative$^{15}$ and $h$ is smaller than a critical field $h_c$, where $\tau_c$, $\tau_w$, and $h_c$ will be determined below. We of course have $u < 0$ if $\bar{u} < 0$ or if the system undergoes a first order phase transition even without the elastic coupling. For $0 \leq h \leq h_c$ the two phases are characterized by $\psi = \psi_0$ and $\psi_1$ with $\psi_1 \geq \psi_0 \geq 0$. As $h \to 0$ we have $\psi_0 \to 0$, while as $h \to h_c$ we have $\psi_1 - \psi_0 \to 0$. We will show that the space average $\langle \psi^2 \rangle$ in the free energy (2.15) gives rise to the two phase coexistence. If it were neglected, we would have the usual tricritical point at $\tau = u = 0$ (see the last paragraph of this subsection$^{15}$).

If the volume fraction of the phase with $\psi = \psi_1$ is written as $\phi$, we have

$$\langle \psi^2 \rangle = \phi\psi_1^2 + (1 - \phi)\psi_0^2.$$  
(3.2)

The average free energy density $\langle f \rangle = F/V$ is given by

$$\langle f \rangle = \phi f_0(\psi_1) + (1 - \phi)f_0(\psi_0) + \frac{1}{4}\beta(\psi_1^2 - \psi_0^2)^2(\phi^2 - \phi).$$  
(3.3)
The interface free energy is neglected. The minimization conditions of $(f)$ with respect to $\psi_1$ and $\psi_0$ are given by

$$f'_0(\psi_1) - \beta(1 - \phi)(\psi_1^2 - \psi_0^2)\psi_1 = 0,$$
$$f'_0(\psi_0) + \beta\phi(\psi_1^2 - \psi_0^2)\psi_0 = 0,$$

which are equivalent to $\delta F/\delta \psi = 0$ at $\psi = \psi_1$ and $\psi_0$. We also minimize $(f)$ with respect to $\phi$ to obtain

$$f_0(\psi_1) - f_0(\psi_0) + \frac{\beta}{4}(\psi_1^2 - \psi_0^2)^2(2\phi - 1) = 0,$$

which means that the two phases have the same free energy density. Note that the quadratic term ($\propto \phi^2$) in $(f)$ in Eq.(3.3) is positive for $\psi_1 - \psi_0 > 0$. Thus, for small $f_0(\psi_1) - f_0(\psi_0)$, a minimum of $(f)$ can be attained as a function of $\phi$ in the range $[0,1]$. These equations may be solved for the simple free energy density (2.2). By eliminating $\phi$ we derive the equations for $\psi_1$ and $\psi_0$ as

$$h/v = \psi_1\psi_0(\psi_0 + \psi_1)^3/3,$$

where $u$ is defined by Eq.(3.1). The negativity of $u$ is required by Eq.(3.8). Thus $\psi_1$ and $\psi_0$ are independent of $\tau$. As $h \to 0$, we have $\psi_0 = 0$ and $\psi_1 = M$, where

$$M = (3|u|/4v)^{1/2}.$$

It is convenient to express $\psi_1$ and $\psi_0$ as

$$\psi_1 = \frac{q}{2} + \sqrt{\frac{q^2}{4} - \frac{3h}{vq^3}}, \quad \psi_0 = \frac{q}{2} - \sqrt{\frac{q^2}{4} - \frac{3h}{vq^3}},$$

where $q$ satisfies

$$h = \frac{2v}{9}q^3(g^2 - M^2).$$

Then $q/M$ is a dimensionless function of $h/vM^5$, tending to unity as $h \to 0$. The difference $\psi_1 - \psi_0 = (q^2 - 12h/vq^3)^{1/2}$ decreases with increasing $h$. A field-induced criticality is attained for $h = h_c$ and $\tau = \tau_c$, where

$$h_c = (8/9)^{5/2}vM^5/12,$$
$$\tau_c = 4vM^4/5 - 2\beta M^2/5.$$  

The critical value of the order parameter is

$$\psi_c = (2/5)^{1/2}M = (3|u|/10v)^{1/2}.$$

For small positive $h_c - h$ we obtain

$$\psi_1 - \psi_0 \approx \frac{2}{9}M(1 - h/h_c)^{1/2}.$$

For $h > h_c$ we have a unique one phase state where $\psi$ is determined by Eq.(2.24). In Fig. 2, we show $\psi_1/M$ and $\psi_0/M$ versus $h/h_c$.

Next the volume fraction of the more ordered phase $\phi$ is calculated. From Eq.(3.5) it depends on $\tau$ as

$$\phi = (\tau_{\text{ex}} - \tau)/\tau_w.$$

This relation holds for $\beta > \bar{u}$ and $h < h_c$ with

$$\tau_{\text{ex}} = -\bar{u}\psi_0^2 - v\psi_1^4 + \frac{v}{3}\psi_1(\psi_0 + \psi_1)^3,$$
$$\tau_w = \beta(\psi_1^2 - \psi_0^2).$$

In Fig. 2, the normalized window width $\tau_w/\beta M^2$ is also displayed as a function of $h/h_c$. Since $\phi$ is in the range $0 < \phi < 1$, the two-phase coexistence is realized in the window region,

$$\tau_{\text{ex}} - \tau_w < \tau < \tau_{\text{ex}}.$$  

For $\tau$ below $\tau_{\text{ex}}$ the more ordered phase starts to appear, and $\tau_w$ is the width of the temperature window. As $h \to 0$, $\tau_{\text{ex}}$ and $\tau_w$ tend to the following values,

$$\lim_{h \to 0} \tau_{\text{ex}} = vM^4/3 = 3u^2/16v,$$
$$\lim_{h \to 0} \tau_w = \beta M^2 = 3\beta(\beta - \bar{u})/4v.$$
On the other hand, as \( h \to h_c \), the upper and lower bounds in Eq.(3.19) meet at \( \tau = \tau_c \) and behave as \( \tau_{cx} \equiv \tau_c + \beta \psi_0 (\psi_1 - \psi_0) \) and \( \tau_{cw} \equiv \tau_c - \beta \psi_0 (\psi_1 - \psi_0) \), where \( \psi_1 - \psi_0 \) depends on \( h_c - h \) as in Eq. (3.15). In Fig. 3, we show the phase diagrams in the \( \tau-h \) plane for \( \bar{u} > 0 \) and for \( \bar{u} < 0 \), separately, where the coexisting curves, \( \tau = \tau_{cx} \) and \( \tau = \tau_{cw} \), and the instability curves are displayed. The latter are obtained by setting \( C\kappa^2 = 0 \) in Eq.(2.22) using \( \psi \) determined by Eq.(2.19) (see the discussions above Eq.(2.23)). These curves meet at the corresponding critical point \( h = h_c \) and \( \tau = \tau_c \) given by Eqs.(3.12) and (3.13).

The usual theory of tricriticality starts with the free energy density,

\[
f = \frac{\tau}{2} \psi^2 + \frac{u}{4} \psi^4 + \frac{v}{6} \psi^6 - h \psi, \tag{3.22}
\]

for systems with short-range interactions. For this model a first order phase transition line appears in the \( \tau-h \) plane for \( u < 0 \). (i) The line starts from the \( \tau \) axis \( (h = 0) \) at the transition point given by \( \tau = 3u^2/16v \) where \( \psi^2 = 3|u|/4v \) in the emerging ordered phase. These values coincide with those in Eqs.(3.20) and (3.9) in our elastic model. (ii) The line ends at a field-induced critical point, where \( \psi^2 = 3|u|/10v, \ h = 8v(3|u|/10v)^{5/2}/3, \) and \( \tau = 9u^2/20v. \) The critical values of \( \psi \) and \( h \) coincide with those in Eqs.(3.14) and (3.12). However, the critical value of \( \tau \) is higher than that in Eq.(3.13) by \( 2\beta M^2/5. \)

B. Magnetization, susceptibility, and specific heat

![FIG. 4: Normalized average magnetization \langle \psi \rangle/M as a function of \( h/h_c \) and \( \tau/\tau_w \) for \( \beta/\bar{u} = 1.2 \) calculated from Eqs.(2.19) and (2.23), where \( M, h_c \), and \( \tau_w \) are defined by Eqs.(3.9), (3.12), and (3.18), respectively.](image)

In the two phase states in the temperature window, the average order parameter is given by

\[
\langle \psi \rangle = \phi \psi_1 + (1 - \phi) \psi_0, \tag{3.23}
\]

which is continuously connected to the solution of Eq.(2.22) in the one phase states outside the window.

![FIG. 5: Normalized susceptibility \( \chi/\chi_c \) as a function of \( h/h_c \) and \( \tau/\tau_w \) for \( \beta/\bar{u} = 1.2 \), where \( \chi_c = M/h_c \). It is calculated from Eqs.(2.20) and (3.24). It increases discontinuously at the phase boundary from the one phase region to the two phase region.](image)

![FIG. 6: Normalized average order parameter \langle \psi \rangle/M, normalized susceptibility \( \chi/\chi_c \), and normalized specific heat \( C_V/C_V_{0} \) versus \( \tau/\tau_w \) for \( \beta/\bar{u} = 1.2 \) in the limit \( h \to 0 \), where \( C_V_{0} = TA^\lambda/2\bar{u} \).](image)

region. See Fig. 4 for \( \langle \psi \rangle \) as a function of \( \tau \) and \( h \) at \( \beta/\bar{u} = 1.2 \). The effective isothermal susceptibility \( \chi = (\partial \langle \psi \rangle/\partial h)_T \) is calculated from

\[
\chi = (\psi_1 - \psi_0) \frac{\partial \phi}{\partial h} + \phi \frac{\partial \psi_1}{\partial h} + (1 - \phi) \frac{\partial \psi_0}{\partial h}. \tag{3.24}
\]

where the derivatives are performed at fixed \( \tau \). See Fig. 5 for \( \chi \) as a function of \( \tau \) and \( h \) at \( \beta/\bar{u} = 1.2 \). We can see that \( \chi \) is discontinuous at the boundary of the window region. There is no critical divergence in \( \chi \) at the field-induced criticality attained. In particular, as \( h \to 0 \), it behaves as

\[
\chi = (1 - 3\phi/4 + 2vM^2/3\beta)/(vM^4/3), \tag{3.25}
\]
where \( vM^4/3 \) is the value of \( \tau_{ex} \) as \( h \to 0 \). For \( \tau > \tau_{ex} \) we have \( \chi = 1/\tau \) at \( h = 0 \). Figure 6 displays the behavior of \( \chi \) on the axis in the limit \( h \to 0 \).

Next we consider the specific heat at constant volume \( C_V = -T\partial^2 f/\partial T^2 \) per unit volume arising from the spin degrees of freedom, where \( h \) is fixed. In the two phase coexistence with \( h < h_c \), we use Eqs.(3.3) and (3.16) to obtain

\[
C_V = TA_0^2/2\beta, \tag{3.26}
\]

which is independent of \( h \) even for \( h > 0 \). In the one phase region, we have \( C_V = TA_0^2\psi^2/(\tau + 3\bar{u}\psi^2 + 5\psi^4) \), where \( \psi \) is determined by Eq.(2.19). In particular, at \( h = 0, C_V = 0 \) for \( \tau > \tau_{ex} \) and \( C_V = TA_0^2/2\bar{u}^2 - 4\bar{u} \) for \( \tau < \tau_{ex} - \tau_w \). In Fig. 6, we show \( C_V \) versus \( \tau \) at \( h = 0 \).

### C. Correlation length and surface tension

![FIG. 7: Inverse correlation lengths \( \kappa_0 \) and \( \kappa_1 \) versus \( h/h_c \) in the coexisting two phases. They are divided by \( \kappa_00 \) in Eq.(3.29). Normalized surface tension \( \gamma/\gamma_0 \) is also shown, where \( \gamma_0 \) is in Eq.(3.36).](image)

Starting with the first line of Eq.(2.27), we may calculate the inverse correlation lengths, \( \kappa_0 \) and \( \kappa_1 \), in the coexisting two phases with \( \psi = \psi_0 \) and \( \psi_1 \), respectively. With the aid of Eqs.(3.6)-(3.8) some calculations yield

\[
\kappa_0^2 = \frac{v}{3C}(\psi_1 - \psi_0)^2(\psi_1 + \psi_0)(\psi_1 + 4\psi_0), \tag{3.27}
\]

\[
\kappa_1^2 = \frac{v}{3C}(\psi_1 - \psi_0)^2(\psi_1 + \psi_0)(4\psi_1 + \psi_0). \tag{3.28}
\]

As \( h \to 0 \), we have \( \kappa_0 \to \kappa_00 \) and \( \kappa_1 \to 2\kappa_00 \), where

\[
\kappa_00 = (v/3C)^{1/2}M^2 \tag{3.29}
\]

is the inverse correlation length in the disordered phase at \( \tau = vM^4/3 \) and \( h = 0 \). As \( h \to h_c \), the inverse correlation lengths go to zero as

\[
\kappa_0 \approx \kappa_1 \approx (4/5)\kappa_00(1 - h/h_c)^{1/2}, \tag{3.30}
\]

from Eq.(3.15). If the scattering amplitude is proportional to \( S_k \) in Eq.(2.21), it grows near the critical point at long wavelengths. In Fig. 7, we plot \( \kappa_0/\kappa_00 \) and \( \kappa_1/\kappa_00 \) versus \( h/h_c \). It is worth noting that the inverse correlation length \( \kappa \) in the one phase region also goes to zero at the criticality. In its vicinity, the relations (2.19) and (2.22) in the one phase case give

\[
C\kappa^2 \approx (h - h_c)/\psi_c, \tag{3.31}
\]

where the term linear in \( \tau - \tau_c \) vanishes.

We also calculate the surface tension \( \gamma \) in the two phase coexistence. We suppose a one-dimensional interface profile \( \psi = \psi(x) \) changing along the \( x \) direction. It changes from \( \psi_0 \) at \( x = -\infty \) and to \( \psi_1 \) at \( x = \infty \). From \( \delta F/\delta \psi = 0 \), we obtain

\[
C\frac{d^2\psi}{dx^2} = f_0'(\psi) - \beta(\psi^2 - \langle \psi^2 \rangle)\psi. \tag{3.32}
\]

We integrate the above equation as

\[
\omega = C(d\psi/dx)^2, \tag{3.33}
\]

where \( \omega(\psi) \) is the grand potential,

\[
\omega = f_0(\psi) - \frac{\beta}{4}(\psi^2 - \langle \psi^2 \rangle)^2 - C_0. \tag{3.34}
\]

From Eq.(3.6) the constant \( C_0 \) in the right hand side can be chosen such that \( \omega \) vanishes at \( x = \pm \infty \) or for both \( \psi = \psi_0 \) and \( \psi_1 \). Some calculations yield

\[
\omega = \frac{v}{3}(\psi - \psi_0)^2(\psi - \psi_1)^2[(\psi + \psi_0 + \psi_1)^2 + \psi_0\psi_1], \tag{3.35}
\]

which turns out to be independent of \( \tau \). The surface tension \( \gamma \) is a function of \( h \) only. It is of the form,

\[
\gamma = \int_{-\infty}^{\infty} dx[\omega + C(d\psi/dx)^2/2] = \int_{\psi_0}^{\psi_1} d\psi \sqrt{2C\omega(\psi)}. \tag{3.36}
\]

In the limit \( h \to 0 \) it becomes

\[
\gamma_0 = \lim_{h \to 0} \gamma = (vC/24)^{1/2}M^4. \tag{3.37}
\]

On the other hand, as \( h \to h_c, \omega \) in (3.35) behaves as \( \omega \approx |u|(\psi - \psi_0)^2(\psi - \psi_1)^2/2 \) so that

\[
\gamma/\gamma_0 \approx (32/375)(1 - h/h_c)^{3/2}, \tag{3.38}
\]

which rapidly decreases near the criticality. See Fig. 7, where \( \gamma/\gamma_0 \) is plotted.
at three regions with $\psi$ measured in units of $M$ and $\kappa$ is in Eq. (4.2) and space and time are measured in units of $L$ and $\ell_0$ in Eq. (4.3). In the initial stage three regions with $\psi \cong \psi_1$ (black), $\psi \cong -\psi_1$ (white), and $\psi \cong 0$ (gray) emerged, but in the final stage $\ell \gtrsim 10^3$ the variant with $\psi \cong -\psi_1$ disappeared here.

IV. NUMERICAL RESULTS

We numerically study the dynamics of our model. We may demonstrate the validity of our equilibrium theory in steady states attained at long times. In our system $\psi$ is a nonconserved variable obeying the relaxation equation,

$$\frac{\partial}{\partial t} \psi = -L_0 \frac{\delta F}{\delta \psi}, \quad (4.1)$$

where $\delta F/\delta \psi$ is given in Eq. (2.14) and $L_0$ is a constant. We integrated the above equation in 2D under the periodic boundary condition. We assume $\bar{u} > 0$ and $\beta/\bar{u} = 1.5$. Then, for $h = 0$, our theory predicts $\psi_1 = 0.612M_0$, $\psi_0 = 0$, $\tau_{\psi_{\infty}}/\tau_0 = 0.047$, $(\tau_{\psi_{\infty}}/\tau_0)/\tau_0 = -0.516$, $\kappa_0\ell = 0.354$, and $\kappa_{\psi}\ell = 0.596$.@ These values will be compared with those from our simulations.

A. Isotropic elasticity

We first assume the isotropic elasticity. We measure $\tau$, $h$, and $\psi$ in units of $\tau_0$, $h_0$, and $M_0$, respectively, where

$$\tau_0 = \bar{u}^2/v, \quad h_0 = v(\bar{u}/v)^{5/2}, \quad M_0 = (\bar{u}/v)^{1/2}. \quad (4.2)$$

FIG. 8: Time evolution of $\psi$ after changing $\tau$ from 0 to $-0.3\tau_0$ at $t = 0$ (upper panel) and steady profile of $\psi/M_0$ obtained at $t = 10^4$ (lower panel) for $\beta/\bar{u} = 1.5$ and $h = 0$ in isotropic elasticity. Here $M_0$ is in Eq. (4.2) and space and time are measured in units of $L$ and $\ell_0$ in Eq. (4.3). In the initial stage three regions with $\psi \cong \psi_1$ (black), $\psi \cong -\psi_1$ (white), and $\psi \cong 0$ (gray) emerged, but in the final stage $\ell \gtrsim 10^3$ the variant with $\psi \cong -\psi_1$ disappeared here.

Here $M/M_0 = [3\beta/\bar{u} - 1]/4]^{1/2}$ from Eq. (3.9). Units of space and time are

$$t_0 = L_0\tau_0, \quad \ell = (C/\tau_0)^{1/2}. \quad (4.3)$$

The scaled time $t_0^{-1}t$ and the scaled space position $\ell^{-1}r$ are simply written as $t$ and $r$ to avoid cumbersome notation. The system size is $200 \times 200$ and the mesh length is $\ell$, so the system length is $200\ell$. In terms of the scaled order parameter $\Psi = \psi/M_0$, Eq. (4.1) is rewritten as

$$\frac{\partial \Psi}{\partial t} = \left[ \nabla^2 - \frac{\tau}{\tau_0} - \Psi^2 + \frac{\beta}{\bar{u}} (\Psi^2 - \langle \Psi^2 \rangle) \right] \Psi + \frac{h}{h_0}, \quad (4.4)$$

As the initial condition at $t = 0$, $\Psi$ at each lattice point consists of a homogeneous constant and a random number in the range $[-0.01, 0.01]$.

In Fig. 8, we show the phase ordering process from a disordered state to a coexisting state. At $t = 0$, $\Psi$ was a random number. For $t > 0$ we lowered $\tau$ from 0 to $-0.3\tau_0$ to induce phase ordering. From our theory, this final $\tau$ is in the coexisting window $[\tau_{\psi_{\infty}} - \tau_{\psi_{\infty}}, \tau_{\psi_{\infty}}]$ and the predicted average order parameter is $0.378M_0$ with $\phi = 0.617$. Since $h = 0$ and $\langle \psi \rangle = 0$ at $t = 0$, the two variants with $\psi = \pm \psi_1$ appeared in the early stage, but the ordered domains with $\psi \cong \pm \psi_1$ disappeared in this run when the domain size became of the order of the system size. (In other runs the variant with $\psi \cong \psi_1$ disappeared as well.) In the steady two phase coexistence at $t = 10^4$ (lower panel in Fig. 8) interfaces are horizontal (parallel to the $x$ axis), where $\psi = 0.612M_0$ in the

FIG. 9: Time evolution of $\psi$ after changing $\tau$ from $-\tau_0$ to $-0.15\tau_0$ at $t = 0$ (upper panel) and steady profile of $\psi/M_0$ obtained at $t = 6 \times 10^3$ (lower panel) for $\beta/\bar{u} = 1.5$ and $h = 0$ in isotropic elasticity. In the phase ordering, ordered regions with $\psi \cong \psi_1$ (black) and disordered regions (gray) emerged. A circular ordered domain remained at long times in this run.
ordered phase and $\langle \psi \rangle = 0.397M_0$. The former coincides with the predicted value, while the latter is slightly larger than predicted.

In Fig. 9, we show the phase ordering process from a one phase state at $\tau = -\tau_0$ to a coexisting state at $\tau = -0.15\tau_0$ at $h = 0$. That is, at $t = 0$, $\Psi$ was the sum of the equilibrium one phase value 0.786 determined by Eq.(2.24) and a random number. The final $\tau$ here is higher than the lower instability value $-0.313^\tau_0$ in Eq.(2.23). Hence phase ordering should take place into a coexisting state where $\phi = 0.350M_0$ and $\langle \psi \rangle = 0.214M_0$ are predicted. In the simulation, regions of the disordered phase appeared, while $\psi$ in the ordered phase changed to $\psi \equiv \psi_1$. In the steady two phase coexistence at $t = 6 \times 10^3$ (lower panel in Fig. 9), a circular ordered domain was realized. There, we find $\psi = 0.594M_0$ in the domain and $\langle \psi \rangle = 0.232M_0$. These values are only slightly different from those predicted.

In Fig. 10, we present a steady profile of $\psi$ at $h = 0.9h_c$ and $\tau = -0.13\tau_0$, where the system is close to the critical point in Eqs.(3.12)-(3.14) and the interface thickness is much widened. For $\beta/\bar{u} = 1.5$, and at this field, our theory gives $\psi_1/M_0 = 0.461, \psi_0/M_0 = 0.304, \tau_{cx}/\tau_0 = -0.032, (\tau_{cx} - \tau_w)/\tau_0 = -0.212, \kappa_0\ell = 0.149$, and $\kappa_1\ell = 0.164$. For the $\tau$ adopted, we predict $\phi = 0.543$ and $\langle \psi \rangle = 0.389M_0$. In the simulation, the maximum and the minimum of $\psi$ are 0.457$M_0$ and 0.302$M_0$, respectively. These values are very close to the theoretical values of $\psi_1$ and $\psi_0$. Furthermore, the observed average $\langle \psi \rangle = 0.392M_0$ is also close to its theoretical average, though the interface regions are very wide here.

### B. Cubic elasticity

Next we integrate Eq.(4.1) in 2D on a cell of $256 \times 256$ assuming the cubic elasticity with $C_{11} - C_{12} = C_{44} = K$.

![Fig. 11: Time evolution of $\psi$ after changing $\tau$ from 0 to $-0.3\tau_0$ at $t = 0$ (upper panel) and final steady profile of $\psi/M_0$ obtained at $t = 10^4$ (lower panel) in cubic elasticity for $\beta/\bar{u} = 1.5$ and $h = 0$. As in Fig. 8, three regions with $\psi \approx \psi_1$ (black), $\psi \approx -\psi_1$ (white), and $\psi \approx 0$ (gray) emerged in the initial stage. Interfaces tend to be parallel to the $x$ or $y$ axis.](image)

where $K = (C_{11} + C_{12})/2$. Then $\xi_a = -1$ from Eq.(2.11) and the softest directions are $[10]$ and $[01]$. As in the isotropic case, space and time are measured in units of $\ell$ and $t_0$ in Eq.(4.3) and we set $\beta = 2\alpha^2/C_{11} = 1.5\bar{u} > 0$. The mesh size of integration is $\ell$. In terms of the scaled order parameter $\Psi = \psi/\psi_0$, the dynamic equation in the 2D cubic case is written as:

$$\frac{\partial \Psi}{\partial t} = \left[ \nabla^2 - \frac{\tau}{\tau_0} - \Psi^2 - \Psi^4 + \frac{\beta}{\bar{u}} G \right] \Psi + \frac{h}{\bar{h}_0}$$

(4.5)

From Eqs.(2.8) and (2.14) we express $G(r)$ in the Fourier expansion,

$$G(r) = \frac{1}{w(0)} \sum_k w(\theta) \Phi_k e^{i k \cdot r},$$

(4.6)

where $\Phi_k$ is the Fourier component of $\Phi = \Psi^2 - \langle \Psi^2 \rangle$ and $w(k) = w(\theta)$ in Eq.(2.13) depends on the angle $\theta$ defined by $\cos \theta = k_x/k$.

In Fig. 11, we lowered $\tau$ from 0 to $-0.3\tau_0$ at $h = 0$ as in Fig. 8. Here the anisotropy of the domain structure arises from the angle dependence of $w(\theta)$ in Eq.(4.6). In the steady state in the lower panel, the maximum of $\psi$ is 0.613$M_0$ and the average $\langle \psi \rangle$ is 0.393$M_0$, in close agreement with the predicted values and those in Fig. 8.

In Fig. 12, we show a steady profile of $\psi$ for $\tau = -0.13\tau_0$ and $h = 0$ as in Fig. 9. Here a square ordered domain is embedded in a disordered region in equilibrium. In the figure, the maximum and the average of $\psi$
field theory. In our model the order parameter $\psi$ is isotropically coupled to the dilation strain $e_1$, as $\psi^2 e_1$, in the free energy, which is the simplest case. Nevertheless, complicated phase behavior follows at constant volume. We summarize our main results.

(i) We have found two phase coexistence in a closed region in the $\tau$-$h$ plane as in Fig. 3. The coexistence region appears under the condition $\bar{u} < \beta$ given in Eq.(3.1). If $\bar{u} > 0$ and $\beta$ is not large, it can be satisfied near the tricritical point. If $\bar{u} < 0$, it can occur even away from the tricritical point.

(ii) The order parameter values in the two phases, $\psi_1$ and $\psi_0$, are determined by $\beta$ only and is independent of $\tau$ as in Fig. 2. The average order parameter $\langle \psi \rangle = \phi \psi_1 + (1 - \phi) \psi_0$ is increased smoothly as $\tau$ is decreased in the window region $\tau_{w} - \tau_{c} < \tau < \tau_{c}$ for $h < h_c$, since the volume fraction $\phi$ depends on $\tau$ as in Eq.(3.16).

The average order parameter $\langle \psi \rangle$ and the susceptibility $\chi = \partial \langle \psi \rangle / \partial h$ are displayed in Figs. 4-6. The specific heat $C_V$ is a constant in two phase coexistence as in Eq.(3.26).

(iii) At the field-induced critical point $h = h_c$ and $\tau = \tau_c$, the correlation length $\xi$ grow and the surface tension $\gamma$ goes to zero as in Fig. 7, while $\chi$ does not diverge.

(iv) We have integrated the dynamic equation, which is Eq.(4.4) for the isotropic elasticity and Eq.(4.5) for the cubic elasticity. A change of $\tau$ from the one phase region into the unstable region induces phase ordering as illustrated in Figs. 8-13. It can occur with decreasing $\tau$ as in Figs. 8 and 11 and with increasing $\tau$ as in Fig. 9. In the final two phase states, the values of $\psi$ and its space average closely agree with the theoretical values.

We make some further remarks.

(i) At constant pressure, two phase coexistence occurs only on a line in the $\tau$-$h$ plane as in the rigid lattice case, but phase separation can be much affected by the elastic coupling (see the appendix). It is worth noting that the transition depends on the sample shape in hydrogen-metal systems at constant pressure, where the proton concentration is linearly coupled to the dilation.

(ii) We mention Monte Carlo simulations on a binary alloy by Landau's group. They assumed that a mixture undergoing unmixing corresponds to ferromagnets and that forming a superstructure to antiferromagnets. In these cases, different results followed in the fixed volume and fixed pressure conditions. However, the unmixing transition in the presence of the size difference is not isomorphic to the ferromagnetic transition. In the former the linear coupling appears between the concentration $c$ and $e_1$ in the form $\psi^2 e_1$, while in the latter the exchange interaction does not break the invariance of $\psi \rightarrow -\psi$ and the elastic coupling is quadratic as $\psi^2 e_1$. At present we cannot compare our theory and their simulations.

(iii) Yamada and Takakura numerically solved a time-dependent Ginzburg-Landau model for an order parameter and a strain in one dimension. They found appearance of a disordered region in a lamellar ordered

V. SUMMARY AND CONCLUDING REMARKS

We have examined the phase transition behavior of compressible Ising models at fixed volume in the mean

FIG. 12: Steady square profile of $\psi/M_0$ in two phase coexistence obtained at $t = 10^4$ in cubic elasticity, where $\beta/\bar{u} = 1.5$, $\tau = -0.15\tau_0$, and $h = 0$.

FIG. 13: Steady, one-dimensional curves of $\psi/M_0$ in two phase coexistence with $\beta/\bar{u} = 1.5$ near the field-induced critical point in cubic elasticity. Here $\tau = -0.13\tau_0$ and $-0.15\tau_0$ for the two curves of $h = 0.9h_c$, while $\tau = -0.13\tau_0$ for $h = 0.95h_c$. The more ordered region expands with lowering $\tau$ at fixed $h$. The values $0.586M_0$ and $0.260M_0$, respectively. The former is slightly smaller than the predicted value $0.612M_0$, while the latter is considerably larger than the predicted value $0.214M_0$.

In Fig. 13, we show one-dimensional steady profiles changing along the $x$ axis near the critical point. The maximum, the minimum, and the average of $\psi$ are $(0.461, 0.305, 0.392)$ for $h/h_c = 0.9$ and $\tau/\tau_0 = -0.13$, $(0.461, 0.308, 0.409)$ for $h/h_c = 0.9$ and $\tau/\tau_0 = -0.15$, and $(0.439, 0.336, 0.398)$ for $h/h_c = 0.95$ and $\tau/\tau_0 = -0.13$. These values closely agree with those from our theory. In these one-dimensional cases, the profiles coincide with those in the isotropic case.
region. Their finding is consistent with our theory. (iii) In real metamagnets, there is no field conjugate to the antiferromagnetic order and the tricriticality has been realized by changing magnetic field or hydrostatic pressure. At fixed volume, our theory predicts two phase coexistence in a temperature window near the tricritical point and near the line of first order phase transition. From Eq.(3.21) the width of the window sensitively depends on the coupling constant $\alpha$ as 

$$\tau_w/\tau_0 = 3\beta(\beta - \bar{u})^2/4\alpha\tau_0,$$

where $\tau_0$ is the coefficient in Eq.(2.3) and $\beta = 2\alpha^2/K$.

(iv) In our mean field theory, we have neglected the renormalization effect near the critical point, which can be intriguing in the presence of the cubic elastic anisotropy. It should be further studied together with the influence of the global elastic constraint studied in this work.

(v) We should generalize our theory to more complex systems. At the ferroelectric transition, the polarization vector is coupled to the strains. In binary alloys, phase separation and an order-disorder phase transition can take place simultaneously, where the concentration $c$ and the structural order parameter $\psi$ are both coupled to $c_1$ in the form $(\alpha_1 c + \alpha_2 \psi^2) c_1$ in the free energy. There can also be a number of anisotropic elastic couplings between the order parameter and the tetragonal or shear strain. We will soon report on phase transition including a Jahn-Teller coupling.

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Appendix A: Fixed pressure condition

We here eliminate the elastic field at fixed pressure. Under isotropic applied stress, we assume an isotropic average dilation change $\langle c_1 \rangle$ caused by the order parameter change. The average stress should be unchanged from that in the reference state, so we require $\langle \sigma_{ij} \rangle = 0$ in Eq.(2.6) to obtain

$$\langle c_1 \rangle = -\alpha\langle \psi^2 \rangle/K,$$  \hspace{1cm} (A.1)

in terms of the bulk modulus $K$. We impose the periodic boundary condition on the deviation, $\delta u_i = u_i - \langle c_1 \rangle x_i/d$, whose Fourier component can be expressed in terms of $\phi_k$ in the same form as that of $u_i$ in the fixed volume case. The free energy consists of $F$ in Eq.(2.15) and

$$\Delta F = -V\alpha^2\langle \psi^2 \rangle^2/2K.$$  \hspace{1cm} (A.2)

The total free energy $F' = F + \Delta F$ is written as

$$F' = \int dr \left[ f + \frac{C}{2}\nabla \psi^2 + \frac{B}{4}\left(\psi^2 - \langle \psi^2 \rangle^2\right)^2 \right],$$  \hspace{1cm} (A.3)

where $f = f_0 - \alpha^2\psi^2/2K$ and $B$ is a positive coefficient,

$$B = 2\alpha^2/K - 2w_M.$$  \hspace{1cm} (A.4)

Here $w_M$ is given by Eq.(2.17). The positivity of $B$ arises from $C_{11} - C_{12} > 0$ and $C_{44} > 0$. The one phase ordered states are determined by $f$. The same form of the free energy was derived by Littlewood and Chandrasekhar for BaTiO$_3$, who argued that the term proportional to $B$ can much decrease the nucleation rate from the paraelectric to ferroelectric state. In our problem, we draw the following conclusion in the mean field theory. In the fixed pressure condition, there can be two phase coexistence only on a first-order coexistence line in the $\tau$-$h$ plane. In fact, $\langle f \rangle$ in Eq.(3.3) would be minimized for $\phi = 0$ or 1 outside the coexistence curve if positive $\beta$ were replaced by negative $-B$.

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For the free energy density (3.22) the first order phase transition line is expressed as
$$\tau = \frac{v (4M^4 - 6M^2 q^2 + 5q^4)}{9}$$
in terms of $q = \psi_1 + \psi_0$, where $q$ is related to $h$ by Eq.(3.11) and $\psi_1 - \psi_0 = \left[(8M^2 - 5q^2)/3\right]^{1/2}$.

At $h = 0$, the two variants with $\psi = \pm \psi_1$ may be present in large systems, while only one of them survived in our simulations in Figs. 8 and 11. Thus we assume $h > 0$ in Eq.(3.23) and take the limit $h \to 0$ in Fig. 6.

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