

Absence of quantum criticality in imbalanced Fermi mixtures at the mean-field level

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1. Introduction

Understanding non-Fermi liquids is among the most interesting challenges to quantum many-body physics [1, 2]. Such systems may occur in a variety of physical contexts such as for example superfluidity in imbalanced Fermi gases. Non-Fermi liquid behaviour could appear in the proximity of a quantum critical point (QCP) [3, 4]. We study the possibility of generating a QCP for imbalanced Fermi gases [5–8] at the mean field level in two and three spatial dimensions. Such systems, containing a mixture of fermionic atoms, nowadays are experimentally available [9–14].

2. Theoretical Model

We study the model defined by the hamiltonian [15]:

$$\hat{H} - \sum_{\sigma} \mu_{\sigma} \hat{n}_{\sigma} = \sum_{\vec{k}, \sigma} \xi_{\vec{k}, \sigma} c_{\vec{k}, \sigma}^{\dagger} c_{\vec{k}, \sigma} + \frac{g}{V} \sum_{\vec{k}, \vec{k}', \vec{q}} c_{\vec{k}+\vec{q}/2, \uparrow}^{\dagger} c_{\vec{k}-\vec{q}/2, \downarrow}^{\dagger} c_{\vec{k}'+\vec{q}/2, \downarrow} c_{\vec{k}'-\vec{q}/2, \uparrow} \quad (1)$$

where $\xi_{\vec{k}, \sigma} = \vec{k}^2/2m_{\sigma} - \mu_{\sigma}$, with $\sigma = \uparrow, \downarrow$, g is an attractive interspecies contact interaction constant and V is the volume of the system. In general, the masses m_{σ} and chemical potentials μ_{σ} can be different.

3. Grand Potential

The grand potential density at the mean-field level can be expressed as:

$$\bar{\Omega} = \frac{\Omega}{V} = \min_{\Delta} \left\{ -\frac{|\Delta|^2}{g} - \frac{1}{\beta} \int \frac{d^d \vec{k}}{(2\pi)^d} \sum_{i \in \{\pm, -\}} \ln(1 + e^{-\beta E_i}) \right\} = \min_{\Delta} \{ \bar{\Omega}_L[\Delta] \}, \quad (2)$$

where \min_{Δ} leads to the global minimum with respect to the order parameter Δ . In this case, there are two possible phases: a normal phase with $\Delta = 0$ and a superfluid phase with $\Delta \neq 0$ [16, 17]. We are denoting the Landau functional as $\bar{\Omega}_L[\Delta]$. Moreover, the elementary excitations energies are given by:

$$E_{\pm} = \frac{\xi_{\vec{k}, \uparrow} - \xi_{\vec{k}, \downarrow}}{2} \pm \sqrt{|\Delta|^2 + \left(\frac{\xi_{\vec{k}, \uparrow} + \xi_{\vec{k}, \downarrow}}{2} \right)^2}. \quad (3)$$

We use $\mu = \frac{\mu_{\uparrow} + \mu_{\downarrow}}{2}$ as the average chemical potential and $h = \frac{\mu_{\uparrow} - \mu_{\downarrow}}{2}$ as the average "Zeeman" field. We are considering a case in which $\mu_{\sigma} > 0$.

4. Landau Coefficients

Setting $\xi = \frac{\xi_{\vec{k}, \uparrow} + \xi_{\vec{k}, \downarrow}}{2}$ we express Landau functional $\bar{\Omega}_L[\Delta]$ as a Taylor expansion in the order parameter Δ :

$$\bar{\Omega}_L[\Delta] = \bar{\Omega}_L(\Delta = 0) + a_2(T)|\Delta|^2 + a_4(T)|\Delta|^4 + O(|\Delta|^6), \quad (4)$$

where the quadratic Landau coefficient a_2 is given by

$$a_2(T) = \left(\frac{\partial^2 \bar{\Omega}_L}{\partial |\Delta|^2} \right)_{|\Delta|^2=0} = -\frac{1}{g} - \frac{1}{4} \int \frac{d^d \vec{k}}{(2\pi)^d} \frac{1}{\xi} \sum_{\sigma} \tanh \left(\frac{\beta \xi_{\vec{k}, \sigma}}{2} \right) \quad (5)$$

and the quartic Landau coefficient a_4 is given by

$$a_4(T) = \frac{1}{2} \left(\frac{\partial^4 \bar{\Omega}_L}{\partial (|\Delta|^2)^2} \right)_{|\Delta|^2=0} = \frac{1}{16} \int \frac{d^d \vec{k}}{(2\pi)^d} \frac{1}{\xi^3} \sum_{\sigma} \left[\tanh \left(\frac{\beta \xi_{\vec{k}, \sigma}}{2} \right) - \frac{\beta \xi_{\vec{k}, \sigma}}{2} \operatorname{sech}^2 \left(\frac{\beta \xi_{\vec{k}, \sigma}}{2} \right) \right]. \quad (6)$$

The QCP correspond to continuous phase transition at zero absolute temperature, so we compute Landau coefficients for $T \rightarrow 0$.

5. Coefficient a_2 for $T = 0$

We use the fact that $\tanh \left(\frac{\beta \xi_{\vec{k}, \sigma}}{2} \right) = 1 - 2f(\xi_{\vec{k}, \sigma})$, where $f(\cdot)$ is the Fermi function, which leads us to the expression below:

$$a_2(T = 0) = -\frac{1}{g} - \frac{1}{2} \int \frac{d^d \vec{k}}{(2\pi)^d} \frac{1}{\xi} \left[1 - \sum_{\sigma} \theta(-\xi_{\vec{k}, \sigma}) \right], \quad (7)$$

where $\theta(\cdot)$ is the Heaviside step function. The first condition of the existence of the QCP requires the disappearance of the coefficient a_2 for that point of the phase diagram.

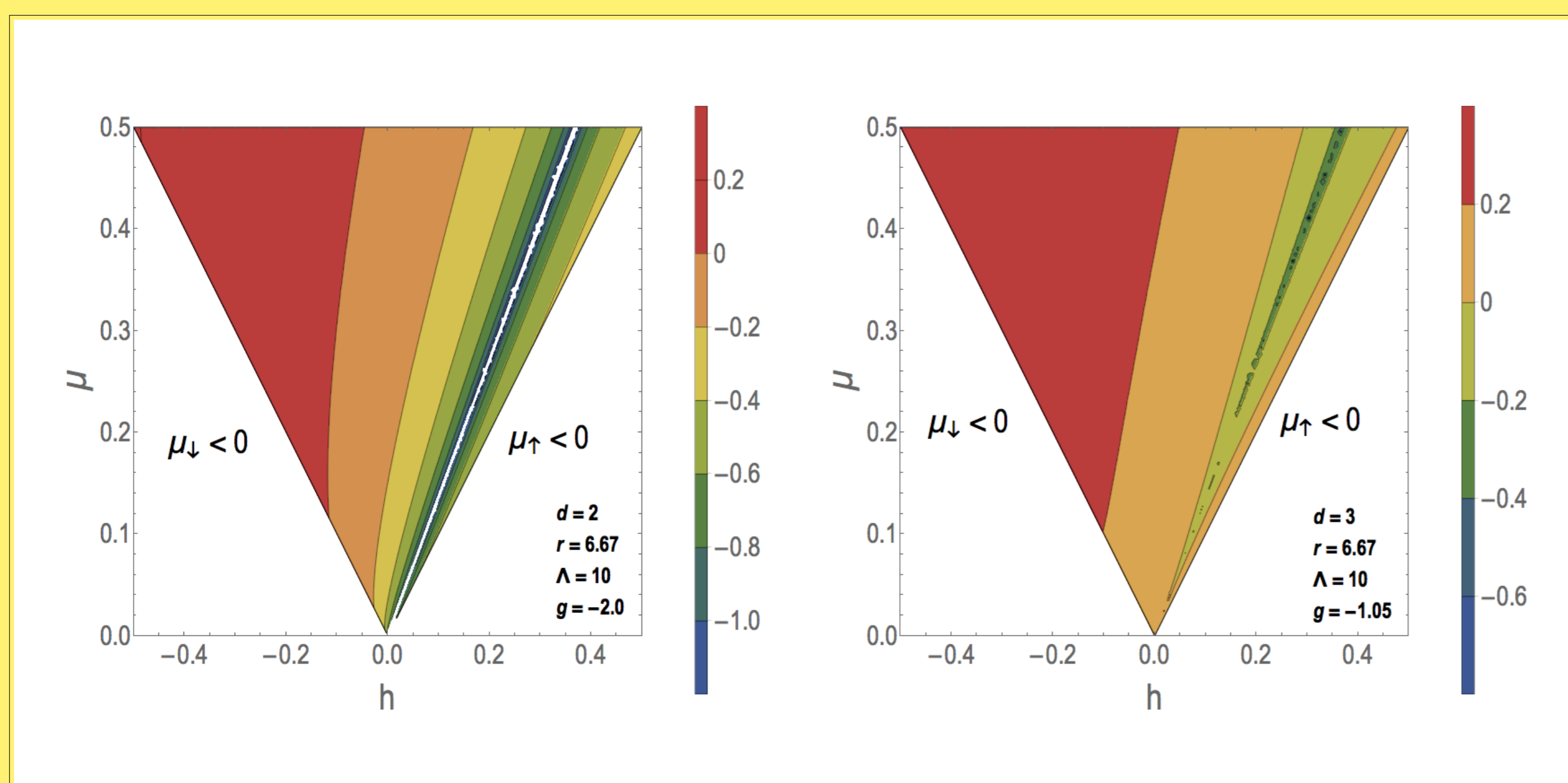


Figure 1: Coefficient a_2 plotted as a function of the Zeeman field (h) and average chemical potential (μ). On the left side we show the plot for $d = 2$, on the right side we show the plot for $d = 3$. In addition, mass ratio r correspond to the case of ${}^6\text{Li}$ and ${}^{40}\text{K}$ mixture.

First we will perform an analysis of this expression in two dimensions:

$$a_2^{2D} = -\frac{1}{g} - \frac{m_r}{4\pi} \ln \left[\frac{|\Lambda^2 - 2\mu m_r| \cdot 2\mu m_r}{|\lambda_{\uparrow}^2 - 2\mu m_r| \cdot |\lambda_{\downarrow}^2 - 2\mu m_r|} \right], \quad (8)$$

where $\frac{2}{m_r} = \frac{1}{m_{\uparrow}} + \frac{1}{m_{\downarrow}}$ is the reduced mass, Λ is the UV cutoff and $\lambda_{\sigma}^2 = 2\mu_{\sigma} m_{\sigma}$. The strength of the interaction can be adjusted using the Feshbach resonance [5–7], so we can always tune the coefficient a_2^{2D} to zero.

Analogously, in three dimensions

$$a_2^{3D} = -\frac{1}{g} - \frac{m_r}{2\pi^2} \left[\Lambda + \frac{\sqrt{2\mu m_r}}{2} \ln \left(\frac{|\Lambda - \sqrt{2\mu m_r}|}{|\Lambda + \sqrt{2\mu m_r}|} \right) - \sum_{\sigma} \left\{ \lambda_{\sigma} + \frac{\sqrt{2\mu m_r}}{2} \ln \left(\frac{|\lambda_{\sigma} - \sqrt{2\mu m_r}|}{|\lambda_{\sigma} + \sqrt{2\mu m_r}|} \right) \right\} \right]. \quad (9)$$

As before, we can tune the a_2^{3D} value to zero. As an example, see figure 1, where we introduce $r = m_{\downarrow}/m_{\uparrow}$.

6. Coefficient a_4 for $T = 0$

Existence of a QCP requires that $a_4 > 0$ in addition to $a_2 = 0$. As before we consider the zero-temperature case:

$$a_4(T = 0) = \frac{1}{8} \int \frac{d^d \vec{k}}{(2\pi)^d} \frac{1}{\xi^3} \left[1 - \sum_{\sigma} \left\{ \theta(-\xi_{\vec{k}, \sigma}) + \xi \delta(-\xi_{\vec{k}, \sigma}) \right\} \right]. \quad (10)$$

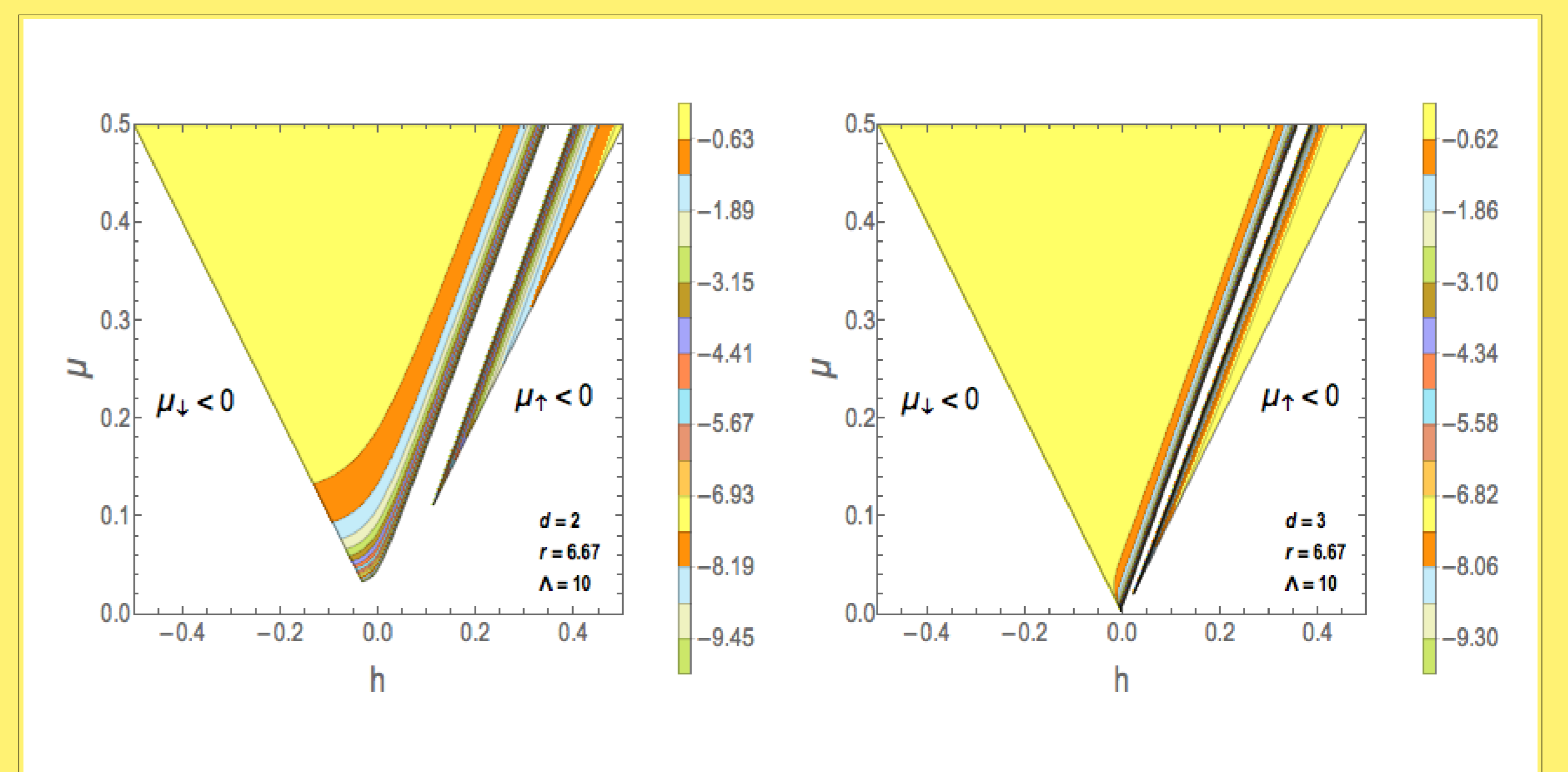


Figure 2: Coefficient a_4 plotted as a function of the Zeeman field (h) and average chemical potential (μ). On the left side we show the plot for $d = 2$, on the right side we show the plot for $d = 3$. Also, in this case, mass ratio corresponds to the lithium and potassium mixture.

In two dimensions we get

$$a_4^{2D} = -\frac{m_r^3}{16\pi} \left[\frac{1}{(\Lambda^2 - 2\mu m_r)^2} + \frac{1}{(2\mu m_r)^2} + \frac{1}{r(\lambda_{\uparrow}^2 - 2\mu m_r)^2} + \frac{r}{(\lambda_{\downarrow}^2 - 2\mu m_r)^2} \right], \quad (11)$$

which is always negative, so in two dimensions there is no QCP at the mean-field level. Furthermore, the coefficient a_4^{3D} in three dimensions is given by

$$a_4^{3D} = -\frac{m_r^2}{64\pi^2} \left[\frac{\Lambda(\Lambda^2 + 2\mu m_r)}{\mu(\Lambda^2 - 2\mu m_r)^2} + \frac{m_r}{(2\mu m_r)^{3/2}} \ln \left(\frac{|\Lambda - \sqrt{2\mu m_r}|}{|\Lambda + \sqrt{2\mu m_r}|} \right) + \sum_{\sigma} \left(\frac{\lambda_{\sigma}(\lambda_{\sigma}^2 + 2\mu m_r)}{\mu(\lambda_{\sigma}^2 - 2\mu m_r)^2} + \frac{m_r}{(2\mu m_r)^{3/2}} \ln \left(\frac{|\lambda_{\sigma} - \sqrt{2\mu m_r}|}{|\lambda_{\sigma} + \sqrt{2\mu m_r}|} \right) - \frac{8\lambda_{\sigma}}{(\lambda_{\sigma}^2 - 2\mu m_r)^2} \right) \right], \quad (12)$$

which as it turns out is also negative, therefore in three dimensions as well there is no QCP at the mean-field level. Figure 2 shows the corresponding graphs.

7. Summary

- We perform a detailed analysis of the conditions necessary for the existence of the QCP in two and three spatial dimensions at the mean-field level.
- In both cases, it appears that no QCP can be generated at the mean-field level.
- Landau coefficients for $T = 0$ are not specified for some parameter choices, such as those corresponding to the BCS theory ($r = 1$, $h = 0$).
- The inclusion of the fluctuations and use of the renormalization group approach allows to modify that result and to generate the QCP in two dimensions [15].

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