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# PHASE TRANSITIONS IN PARTIALLY SPIN-POLARIZED <sup>3</sup> He

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#### <u>Résumé</u>

Nous présentons les diagrammes d'équilibre liquide-gaz pour l'hélium trois partiellement polarisé. A basses températures, nous trouvons des phénomènes intéressants tels que le métamagnétisme en phase liquide ou une surpolarisation du liquide.

#### Abstract

We present liquid-gas equilibrium phase diagrams for partially spin-polarized <sup>3</sup>He. At low temperatures, interesting phenomena such as liquid metamagnetism and liquid overpolarization have been found.

# Introduction

The study of phase transitions in spin polarized <sup>3</sup>He is a subject of growing interest. At high pressures, the results so far obtained (1) might indicate the existence of a metamagnetic, or near it, transition in the liquid phase. In this contribution, we want to investigate the properties of polarized <sup>3</sup>He at lower pressures, in particular at the vapor pressure. This study has been prompted by some experiments on the liquid-gas phase transition which are now in progress at the Ecole Normale de Paris (2).

To investigate liquid-gas (and liquid-liquid) phase transitions, we resort to a simple model already employed in Nuclear Physics (3). It is based on the use of Skyrme-like forces in the framework of a mean field approach (4). A more detailed description of the method as well as further results can be found in refs 5 and 6.

# The formalism

A convenient thermodynamic potential to describe the system is the free energy density that per unit volume reads

$$f(\boldsymbol{g}, \boldsymbol{M}, \boldsymbol{T}) = f_{n,i}(\boldsymbol{\varrho}, \boldsymbol{M}, \boldsymbol{T}) + f_{v}(\boldsymbol{\varrho}, \boldsymbol{M}, \boldsymbol{T})$$
(1)

where g is the number density, g = N/V, T is the temperature and M the magnetization

 $M = (g^{\uparrow} - g^{\downarrow})/\rho$  (2)

**g**<sup>1</sup> and **g**<sup>1</sup> are the spin up (down) partial densities. In expression 1,  $f_{n,i}$  is the free energy density of a non-interacting Fermi gas (7) and  $f_{v}$  is the contribution due to the interactions. This is the key ingredient for which we have taken the following function:

$$f = 1/2(b_0 + b_1 M^2 + b_2 M^4) g^2 + (3)$$

$$1/2(c_0 + c_1 M^2 + c_2 M^4) g^{2+6}$$

To fix the coefficients entering eq. 3 we have used as much experimental information as possible, which is mainly available for M around 0.

First of all, the values of  $b_0$ ,  $c_0$  and  $\sigma$  at T=0 were fixed in order to reproduce the experimental values of the energy per particle  $E_0$ , equilibrium density  $g_0$  and compressibility  $\mathbf{K}$  of unpolarized liquid <sup>3</sup>He (5). Next, the coefficient  $b_0$  was made T-dependent and fixed to exactly reproduce the vapor pressure of unpolarized <sup>3</sup>He. To obtain  $b_1(T)$  and  $c_1(T)$ , we have used the magnetic susceptibility  $\mathbf{K}$ 

$$\frac{c}{\chi} = \frac{1}{g^2} \frac{\partial^2 f}{\partial M^2} \Big|_{T}$$
<sup>(4)</sup>

for which there are experimental results at M=0. In eq. 4, C is the Curie's constant.

The necessity of at least M<sup>4</sup> terms in eq. 3 stems from experimental facts and theoretical arguments. Indeed, at M $\sim$ 0 it is known experimentally that it is easier to magnetize the vapor than the liquid. Consequently, in the liquid-gas equilibrium, near M=0 the phase diagram would be that of fig. 1, where we show the pressure P as a function of M for a given T (L reads for the liquid, G for the gas in equilibrium with it). If P(M) were always an increasing function of M, as it is the case of fig. 1, eq. 3 <u>without</u> M<sup>4</sup> terms would be fine. However, it is easy to show that this is not so for all T.

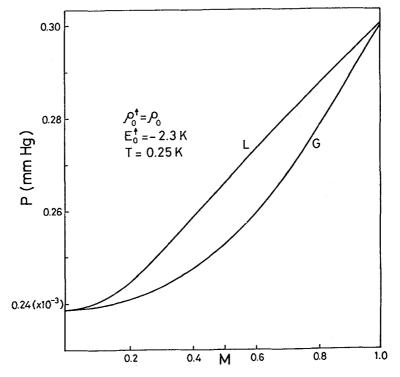


Figure 1

To see how it comes about, let us consider a very low T and write the pressures  $P(P^{\uparrow})$  of the M=O(1) vapor as a function of the energy per particle  $E_o(E_o^{\uparrow})$ :

$$P = A T^{5/2} exp(E_o/T)$$

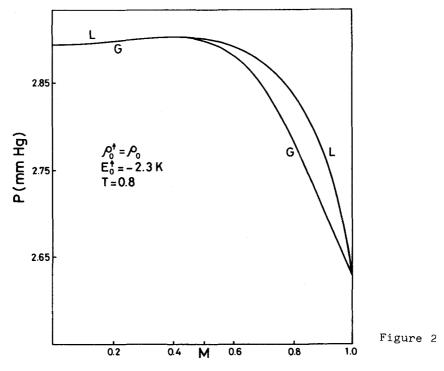
$$Pt = 2 A T^{5/2} exp(E_ot/T)$$
(5)

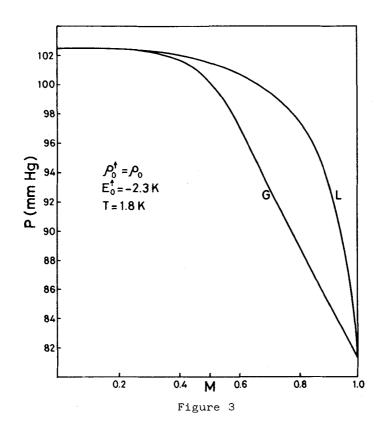
where A is a constant. Since  $|E_o\uparrow| < |E_o|$ , at T low enough we have P↑>P. But when T increases, there is a temperature T\* for which both pressures are the same

$$T^* = (E_0^{\dagger} - E_0) / \ln 2$$
  $\Delta E / \ln 2$  (6)

and from T\* up, P> P1. Recent calculations (8) show that  $E_0 \sim E_0^{\uparrow}$  and  $g_0^{\uparrow} \sim g_0^{\circ}$ . That makes T\* rather small. For example, if  $\Delta E=0.2$  K, T\*  $\sim 0.3$  K, much smaller than the critical temperature of <sup>3</sup>He (5) (T\* estimated from eq. 6 can be accurate only if T\* is not very high. For  $\Delta E= 0.2$  K our calculations yield T\*  $\sim 0.5$  instead of 0.3 K). We thus conclude that in order to bend down the P(M) curve, we do need M<sup>4</sup> terms. Finally, the values of  $b_2(T)$  and  $c_2(T)$  are essentially fixed by the values of the experimentally unknown quantities  $E_0^{\uparrow}$  and  $g_0^{\uparrow}$ (6) which are the free parameters of our model.

The presence of  $M^4$  terms in eq. 3 makes the phenomenom of liquid metamagnetism (divergence of  $\chi(M)$  for some values of M) possible. For given values of g and T, the function  $\chi(M)$  becomes negative in some range of values of M. Regions of negative  $\chi$  or negative compressibility K are instability regions and the system has to separate in several phases that differ in density and/or magnetization.





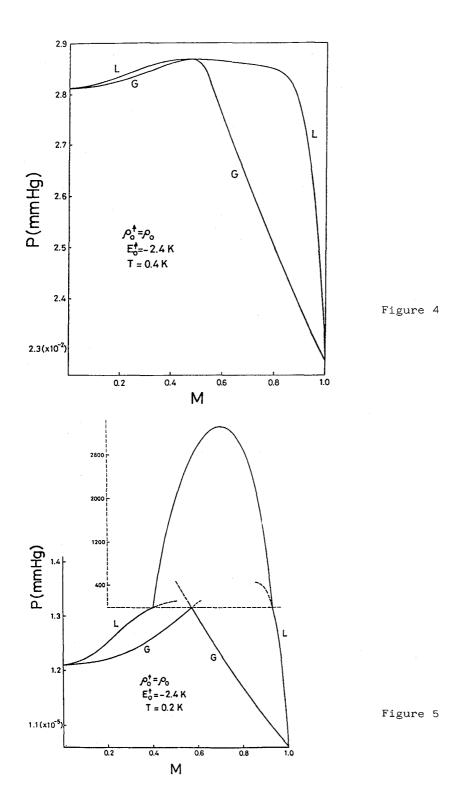
Results

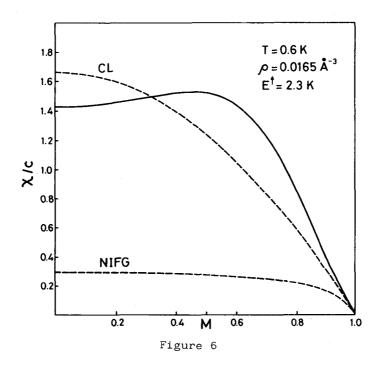
Typical phase diagrams for selected values of T are shown in figs. 1-5. Figures 1 to 3 correspond to  $E_0 = -2.3$  K (this energy is the more recent theoretical estimate, see ref. 8; the results are rather insensitive to the value of  $g_0$  which is also estimated in that ref. to be smaller than  $g_0$ ), whereas figs. 4 and 5 correspond to  $E_0 = -2.4$  K.

In fig. 1 we present the situation  $T \leq T^*$  whereas figs. 2 and 3 correspond to  $T > T^*$ . One may see in fig. 3 that for  $M \ge 0.4$  the liquid is more magnetized than the vapor, in contradistinction with fig. 1. The same phenomenon of liquid overpolarization is presented in fig. 4 (T\* for  $E_0^{T} = -2.4$  K is  $\sim 0.2$  K).

Figure 5 is the result of combining magnetic (  $\chi < 0$ ) and mechanical ( $\chi < 0$ ) instabilities. At low pressures we have a liquid -gas equilibrium that shows again liquid overpolarization. At high pressures, two liquids with rather different magnetizations and similar densities are in thermodynamical equilibrium. When P decreases, there is a point where two liquids are in equilibrium with a <u>common</u> vapor of intermediate M (triple line). We have not found a quadruple point in the calculations performed with our model functional, eq. 3.

Finally, figure 6 shows  $\chi(M)$  for  $E_0^{\dagger} = -2.3$  K at T= 0.6 K. In this figure, the curve labeled CL corresponds to the classical gas, which is a good approximation for the vapor at this temperature. NIFG represents the non-interacting Fermi gas susceptibility and the solid line is  $\chi(M)$  calculated from eqs. 1, 3 and 4. Notice that near M=0, the magnetic susceptibility of the gas is larger than that of the liquid, whereas near M=1 the opposite occurs. This is so because T > T\* and is in agreement with the situation shown in figures 2 and 3.





# <u>Conclusions</u>

Our simple model predicts interesting physical situations and shows the key role played by  $E_0$  in the thermodynamics of partially polarized <sup>3</sup>He. Our predictions can be of some relevance for future experiments. We have found that one has to go beyond M $\sim$  0.5 to find the most interesting effects.

Even if metamagnetism does not show up, situations near it might well be met (figs. 2 and 3), playing an important role (9). One could imagine purely thermal overpolarization methods taking advantage of the particular shape of these diagrams.

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