Polarized $^3$He as a probe for short range spin-dependent interactions

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(Dated: October 20, 2010)

We have studied the relaxation of a spin-polarized gas in a magnetic field, in the presence of short-range spin-dependent interactions. As a main result we have established a link between the specific properties of the interaction and the dependence of the spin-relaxation rate on the magnitude of the holding magnetic field. This allows us to formulate a new, extremely sensitive method to study short-range interactions which yields a two-order-of-magnitude improved constraint on the coupling strength $(g_s g_p)$ as a function of the force range $(\lambda)$: $g_s g_p \lambda^2 < 3 \times 10^{-27}$ m$^2$.

PACS numbers: 14.80.Va, 67.30.ep

Hyperpolarized $^3$He is currently applied to a wide variety of scientific and medical problems. They include magnetic resonance imaging, spin-polarized targets, surface science, probing of biological systems, and precision measurements in fundamental physics \[1\] \[14\]. One of the unique properties of polarized $^3$He is the very long spin-relaxation time constant which can be of the order of thousands of hours \[11\] \[12\], making polarized $^3$He extremely sensitive to any spin-dependent interaction. It is well known that the presence of a magnetic field gradient in a cell containing spin-polarized gas significantly affects the spin-relaxation. The origin of this relaxation mechanism is the loss of phase coherence of the atoms due to the fluctuating magnetic field seen by the atoms as they diffuse throughout the cell. Spin relaxation of a gas in the presence of a uniform gradient has been analyzed by Cates, Schaefer and Happer using a perturbation theory approach \[13\] and by McGregor \[14\] within the framework of Redfield theory. In the present work we focus on spin-relaxation phenomena due to the field gradient decaying over distances much shorter than the cell size. This results in new expressions for the relaxation rates which agree with the already known result \[13\] \[14\] in the limit of high pressure and high magnetic field (adiabatic regime) and in the uniform gradient limit for very low pressure and magnetic field (“motion narrowing” regime). A broad transitional regime of spin motion has been discovered which carries the complete information on the strength of this short-range field as well as on its spatial dependence. This finding allows us to propose a new very sensitive method to study short-range spin-dependent interactions from various origins, e.g. the magnetic field of a strongly diluted ferromagnetic sample or the pseudomagnetic field of hypothetical axionlike forces \[15\] \[17\]. To test the power of the method we have performed an experiment with polarized $^3$He measuring the longitudinal relaxation rate as a function of applied magnetic field. The experiment results in a new, stronger constraint on the axionlike interactions.

Suppose the magnetic field in a cell may be described by a homogeneous magnetic field $B_0$ and weak inhomogeneous field $b(r)$: $B(r) = B_0 + b(r)$, with $< b(r) > = 0$. According to Slichter \[18\], from the Redfield theory of spin-relaxation due to a randomly-fluctuating magnetic field, the spin-relaxation rates are given by

\[
\Gamma_1 = \frac{1}{T_1} = \frac{\gamma^2}{2} [S_z(\omega) + S_y(\omega)],
\]

\[
\Gamma_2 = \frac{1}{T_2} = \frac{\gamma^2}{4} [2S_z(\omega) + 2S_y(\omega) + 2S_z(0)].
\]

Here $T_1$ is the time constant for the longitudinal relaxation, $T_2$ is the time constant for the transversal relaxation, $\gamma$ is the gyromagnetic ratio for the atoms of the gas ($\gamma \approx 2.04 \times 10^4$ s$^{-1}$ G$^{-1}$ for $^3$He), and $\omega = \gamma B_0$ is the Larmor frequency. The functions $S_{k=x,y,z}(\omega)$ are the Fourier transform components of the magnetic field autocorrelation function $R_k(\tau) = \langle b_k(t)b_k(t + \tau) \rangle$, where the ensemble average can be evaluated as

\[
R_k(t - t_0) = \int \int \rho(r_0, t_0)\rho(r, t|r_0, t_0)b_k(r)b_k(r_0)d^3r_0 d^3r, \tag{3}
\]

knowing the conditional probability density $\rho(r, t|r_0, t_0)$ for an atom sitting at time $t_0$ at $r_0$ to be found at later time $t$ at a position $r$. The factor $\rho(r_0, t_0)$ is the single probability density. For times $|t - t_0|$ much longer than the mean time between atomic collisions, the conditional density obeys the diffusion equation \[19\], with the constraint that the initial $\rho(r, t_0|r_0, t_0) = \delta(r - r_0)$ and reflection boundary conditions. Let us consider a rectangular cell of length $L$ with square base of size $R$. Let $x$ be the cell axis so that the square ends occur at $x = \pm L/2$, a homogeneous field $B_0$ directed along $z$ axis, and an inhomogeneous magnetic field $b(x)$ directed along $x$ axis. For this geometry the problem becomes unidimensional with a known analytical solution for $\rho$ \[14\]. Substituting this solution in (3) and taking into account that in our case $\rho(r_0, t_0) = 1/L$ we get the following expression for...
the autocorrelation:

$$R_z(\tau) = 2 \sum_{n=0}^{+\infty} \frac{1}{\tau_n} \frac{1}{\pi^2(2n+1)^2} b_{x,n}^2$$  \hspace{1cm} (4)

$$\tau_n = \frac{\tau_L}{\pi^2(2n+1)^2}$$

$$b_{x,n} = \int_{-L/2}^{L/2} b(x) \sin((2n+1)\pi x/L) \frac{dx}{L}$$  \hspace{1cm} (5)

where we have introduced the characteristic diffusion time constant $\tau_L = L^2/D$, $D$ is the diffusion coefficient of the gas. Taking the Fourier transform gives

$$\Gamma_1(\omega) = \frac{\gamma^2}{2} S_x(\omega) = 2 \gamma^2 \sum_{n=0}^{+\infty} \frac{1}{\tau_n} \frac{1}{1+(\omega \tau_n)^2} b_{x,n}^2$$  \hspace{1cm} (6)

To proceed further we have to specify the field $b(x)$. To take a concrete example, we consider the macroscopic pseudomagnetic field representing an axionlike interaction of polarized $^3$He with the nucleons in the cell walls [20]:

$$b(x) = \frac{h}{2\gamma m_n} N g_p g_p \left(1 - e^{-d/\lambda}\right),$$  \hspace{1cm} (7)

$$b_a = \frac{h\lambda}{2\gamma m_n} N g_p g_p \left(1 - e^{-d/\lambda}\right).$$  \hspace{1cm} (8)

Here, $x$ is the distance from the wall, $g_s$ and $g_p$ are dimensionless scalar and pseudoscalar couplings between the nucleon and the axionlike particle, $\lambda = \frac{h}{m_n}$ is the force range, $m_n$ is the nucleon mass, $N$ is the nucleon number density and $d$ is the thickness of the walls. Substituting (7) in (5)-(6) we arrive at a general expression for the longitudinal relaxation on the one-dimensional case:

$$\Gamma_1 = (\gamma b_a)^2 \tau_\lambda \left(1 + e^{-L/\lambda}\right)^2 \left(\frac{\cos \sqrt{\phi_L/2} + \cosh \sqrt{\phi_L/2}}{\sinh \sqrt{\phi_L/2}}\right)^{-1} \left(1 + \phi_\lambda^2 + \frac{\lambda}{2}(\phi_\lambda^2 - 3) \sinh \frac{L}{2\lambda}\right)$$  \hspace{1cm} (9)

![FIG. 1: Relaxation rate due to a short-range gradient magnetic field $b$ versus the magnitude of the homogeneous magnetic field calculated according to (9) for three different values for the range $\lambda$.](image)

where $\phi_L = \omega \tau_L$, $\phi_\lambda = \omega \tau_\lambda$ and $\tau_\lambda = \lambda^2/D$.

Our result (9) is illustrated in Fig. 1. We can distinguish three regimes of relaxation: relaxation in a low magnetic field $\omega \ll \tau_L^{-1}$ with $\Gamma_1 \propto const$, relaxation in a moderate field $\tau_L^{-1} \ll \omega \ll \tau_\lambda^{-1}$ with $\Gamma_1 \propto \omega^{-1/2}$, and relaxation in a high field $\omega \gg \tau_\lambda^{-1}$ with $\Gamma_1 \propto \omega^{-2}$.

The magnetic field value corresponding to the transition between the latter two regimes depends on the force range $\lambda$: the smaller $\lambda$ is, the higher magnetic field is needed. For an extremely sharp correlation function...
tions with the cell walls, all components of the pseudomagnetic field $b_r(x)$, $b_y(y)$, and $b_z(z)$ need to be taken into account. Since the Brownian motions of the three coordinate directions are independent, from (11) follows

$$\Gamma_1 = \Gamma_{1,x} + \Gamma_{1,y}$$

where both $\Gamma_{1,x}$ and $\Gamma_{1,y}$ are given by our expression (13) for the one-dimensional geometry. The transverse relaxation rate follows from eqns. (2) and reads, for a rectangular (or cubic) cell

$$\Gamma_2 = \Gamma_1(0) = 2(\gamma b_0)^2 \tau_\lambda \quad \text{for} \quad \lambda \ll R, L.$$  

(13)

We expect similar result for other geometries (cylindrical, or spherical) since (13) is independent of the cell size. We checked by Monte Carlo simulation that this is indeed the case at the 10% level of precision.

We now apply our theoretical results to search for an exotic short-range axionlike interaction. Assuming for the time being no extra interactions, the experimental spin-relaxation is determined by contributions from three sources: $\Gamma_{exp} = \Gamma_{dd} + \Gamma_{wall} + \Gamma_m$, where $\Gamma_{dd}$ is the dipole-dipole relaxation due to atomic collisions, $\Gamma_{wall}$ is due to the $^3$He spin relaxation on the walls of the cell and $\Gamma_m$ is due to magnetic field inhomogeneities. $\Gamma_{dd}$ and $\Gamma_{wall}$ are expected to be independent of magnetic field. If $B_0$ is high, so that $\omega \gg \tau_L^{-1}$, the relaxation due to magnetic field inhomogeneities can be decomposed into two components as follows:

$$\Gamma_m \approx D \left( \langle |g_e|^2 \rangle + \langle |g_i|^2 \rangle \right) / B_0^2 = \Gamma_{mi} + \Gamma_{me}$$  

(14)

where $g_i$ is the gradient due to inhomogeneities of the holding magnetic field $B_0$ and $g_e$ is the gradient caused by the external magnetic environment. Since $g_i \propto B_0$, the relaxation $\Gamma_{mi}$ due to the magnetic field gradient caused by the environment is the only term that depends on $B_0$. As all external magnetic sources are far away from the cell containing the polarized $^3$He, the relaxation is only affected by magnetic sources which are large compared to the cell size, and hence providing a nearly uniform gradient over the cell volume. Thus, we can expect the $\Gamma_{me}$ scales as $B_0^{-2}$. Finally, we can write:

$$\Gamma_{exp}(B_0) = \Gamma_{dd,wall,mi} + D|g_e|^2 B_0^{-2}.$$  

(15)

The relaxation due to short-range spin-dependent forces \([9]\) depends very differently on the magnetic field compared to the simple law \([15]\), and can thus be separated from the other sources of relaxation. To take advantage of this feature we performed measurements of the longitudinal relaxation of $^3$He as a function of magnetic field using a cylindrical alumino-silicate glass cell (GE180, 5 cm diameter, 10 cm long, 3 mm wall thickness) filled with polarized $^3$He gas (75% initial polarization at 0.3 bars pressure) placed inside a self-screening “Magic box” magnetostatic cavity \([24]\). More details of the experiment will be published elsewhere. The experimental results obtained are shown in Fig. 2 together with a fit of expression (15):

$$\Gamma_{dd,wall,mi} = (9.71 \pm 0.16) \times 10^{-3} \text{ h}^{-1},$$

$$|g_e| = (2.70 \pm 0.05) \times 10^{-2} \text{ G/m}.$$  

(16)

FIG. 2: Experimentally measured longitudinal relaxation rate of polarized $^3$He versus magnetic field (points). Solid line shows a fit of the theoretical prediction (15) for no short-range spin-dependent forces.

Now we consider the addition of an axion rate \([9]\) to the normal expression \([15]\). The fit involves now two additional free parameters $\lambda$ and $b_0$, it yields a result compatible with zero for $b_0$. Thus the experimental data shows no evidence for a new axionlike interaction. In order to set an upper limit on the strength of an axionlike interaction, the fit was performed again with various fixed values of $\lambda$ ranging from one micron to one centimeter. For each such value of $\lambda$ an upper limit on the product $g_3q_3$ was derived from the fit, which is shown as bold dashed line in Fig. 3. A previous attempt \([25]\) to determine constraints on axionlike forces in the range $10^{-6} \text{ m} < \lambda < 10^{-2} \text{ m}$ using available $^3$He relaxation data \([11, 12]\) appears to be flawed. That analysis was based on expression (12), which is only valid in the limit of high pressure and high magnetic field: $\omega \gg \tau_\lambda^{-1} \gg \tau_L^{-1}$, which corresponds to $\lambda > 10^{-4} \text{ m}$ for the experimental conditions \([11, 12]\).

During the time that the present studies were performed, new experimental data on the very long time constant of the transverse relaxation in the “motion narrowing” regime became available from the shielded room BMSR-2 in Berlin \([26]\). We have performed an analysis of these data within our theory. For the experimental conditions of \([26]\) the transversal relaxation time constant may be written as follows \([13, 14]\):

$$\frac{1}{T_2} = \frac{1}{T_1} + \frac{4\gamma^2 R^4}{175D} (\nabla B_x^2 + \nabla B_y^2 + 2\nabla B_z^2)$$

(17)

where the first term represents the field-independent longitudinal relaxation due to interaction with walls and atomic collisions, while the second term represents the relaxation due to long-range gradients of the magnetic field. The first term \([17]\) was measured in a high magnetic field: $T_1 = 85 \pm 5 \text{ h}$. The second term was estimated to be $T_{2m} = 370 \pm 64 \text{ h}$ from the measured values of the gradients in \([17]\). The authors \([26]\) compared the sum...
formulated, supported by a first experiment: scanning the relaxation rate $\Gamma_1$ on the holding magnetic field $B_0$. In general, this method samples the inhomogeneous field $b(x)$ at different spatial scales. Information about $b(x)$ can be extracted from comparison of the data $\Gamma_1(\omega)$ with the expectations of the model. In the particular case of the simple parametrization (7), one can extract the amplitude $b_0$ and the range parameter $\lambda$. Our first experiment could be dramatically improved if performed in a “zero field room” such as BMSR-2, where the external gradient of the magnetic field is 4 orders of magnitude lower than in our “Magic box”.

We are grateful to V. Nesvizhesky for attracting our attention to the axion problem and to E. Kats and R. Whitney for valuable discussions. One of us (A. P.) expresses his gratitude to C. Fu and T. Gentile for fruitful discussion during a visit at NIST.

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