

Low field depolarization of electronic spins through dipole-dipole coupling

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C'est trop bien

Blabla sur les NV, les ensembles etc.

In this letter/article, we characterize the depolarization of the spins observed for dense ensemble of NV centers in zero magnetic field and its potential application for DC magnetometry. While the main mechanism behind the depolarization, the lift in the degeneracy between the four classes of NV centers, is already well studied and can be exploited in a microwave-less vector magnetometry protocol, we found two other depolarization mechanisms specific to the zero-field region which could play an important role in the low-field magnetometry protocol.

The T_1 depolarization dynamics of single or sparse NV centers at room temperature is dominated by a two-phonon Raman process, which depends on the crystal lattice temperature but does not (?) depend on the external magnetic field. However, it has been observed that dense NV centers ensemble have an additional spin lifetime contribution which depends greatly on the magnetic field and not on the temperature. This effect has been attributed to cross-relaxation between the NV centers through dipole-dipole coupling. Some inhomogeneity between the NV centers is further needed in order to explain the depolarization of the spin ensemble. We will denote T_1^{ph} the characteristic time associated with the phonon relaxation process and T_1^{dd} the characteristic time associated with the dipole-dipole relaxation process.

The main signature of the spin depolarization in low field is the characteristic dip in photoluminescence (PL) observed only for high density ($\gtrsim 1$ ppm) of NV centers, as shown on fig 1. The decrease in the spins' lifetime in zero field makes the optical polarization scheme of the NV centers less effective and therefore reduce the population of the bright $|0\rangle$ spin state. The reason for the

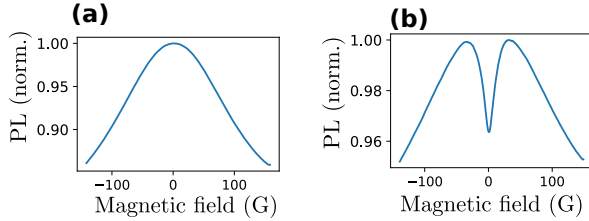


FIG. 1. Photoluminescence from NV centers as a function of an external magnetic field randomly oriented for two samples : (a) a CVD sample containing ≈ 4 ppb NV^- centers, (b) an HPHT sample containing ≈ 3 ppm NV^- centers

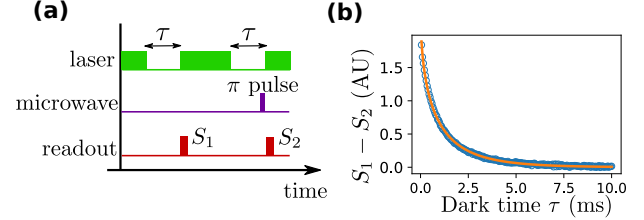


FIG. 2. Spin T_1 measurement protocol. (a) Sequence used to measure the spin lifetime : the laser is turned off for a time τ twice and the spin state is read at the end of the dark time by measuring the initial photoluminescence when the laser is turned back on. A microwave resonant with one of the spin transitions is used in one of the sequence to project the $|0\rangle$ spin state in either $|+1\rangle$ or $|-1\rangle$. The sequence result is the subtraction $S_1 - S_2$. (b) Measurement of an ensemble spin lifetime, fitted with an expression $A \exp\left(-\frac{\tau}{T_1^{\text{ph}}} - \sqrt{\frac{\tau}{T_1^{\text{dd}}}}\right)$

decrease in PL at higher magnetic field values is the mixing of the bright spin state $|0\rangle$ to the darker state $|-1\rangle$ induced by the transverse magnetic field. This effect does not modify (on first approximation) the spins' lifetime, and is common to both dense and sparse samples.

The spin lifetime protocol used here is described in Fig. 2 and is based on previous similar experiments. It consists in a pump-probe measurement where the spins are first polarized in the $|0\rangle$ state by a green laser, and read-out optically after a variable dark time τ . Using only this sequence can result in artifacts in the signal, mostly due to charge state transfer in the dark [2]. It is therefore convenient to repeat the sequence with an additional π pulse right before the spin read-out to project the remaining $|0\rangle$ polarization into a darker $|+1\rangle$ or $|-1\rangle$ state. By subtracting the result of the two sequences, we select only the spin-dependent part of the signal, with the added benefit of being able to select a specific class of NV centers.

The ensemble spin lifetime is non-exponential due to inhomogeneities between the NV^- centers. We will base our interpretations of the experimental results on the NV-fluctuator model developed in [1]. A conclusion of this model is that, for an homogeneous 3D distribution of fluctuator, the dipole-induced lifetime should be stretched-exponential with a stretch factor $\beta = 1/2$. Such an ensemble lifetime is indeed observed when $T_1^{\text{dd}} \ll T_1^{\text{ph}}$ (see SI).

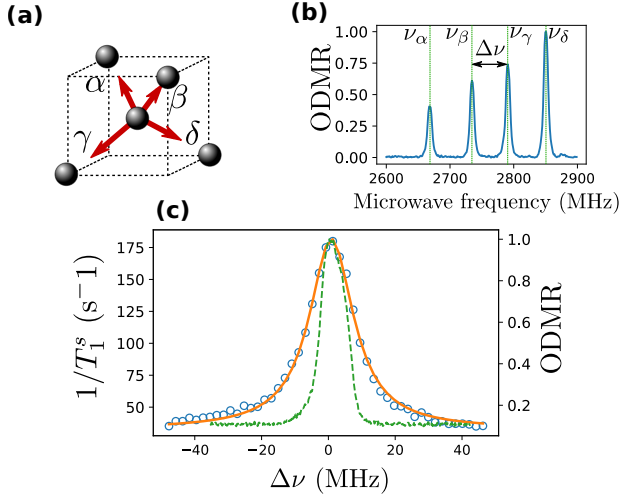


FIG. 3. Tuning resonant coupling through magnetic field orientation. (a) Sketch of the four possible orientations ("classes") in a single crystal diamond lattice. (b) ODMR spectrum showing four $|0\rangle \rightarrow |-1\rangle$ resonances corresponding to the four spin classes. The detuning $\Delta\nu$ between the classes β and γ was controlled by changing the orientation of the external magnetic field. (c) Stretch part of the lifetime decay for the spins resonant with ν_γ as a function of the detuning $\Delta\nu$ (blue circles), fitted by a Lorentzian with half width at half maximum 8.04 MHz. Green dashed line correspond to the ODMR width of a single class stretched by a factor $\sqrt{2}$, approximating the overlap of the two classes (see SI).

For the samples used in this paper (detailed in SI), we observed that $T_1^{\text{dd}} \sim T_1^{\text{ph}}$, meaning that we had to include both in our analysis. The resulting fitting formula is therefore :

$$S(\tau) = A \exp\left(-\frac{\tau}{T_1^{\text{ph}}} - \sqrt{\frac{\tau}{T_1^{\text{dd}}}}\right), \quad (1)$$

In order to get comparable fitting parameters between each measurement, we decided to fix the value of the exponential lifetime in the fitting process, leaving only the stretch lifetime as a free parameter. For Fig. 2, 3, 4 and 6, the chosen exponential lifetime was $T_1^{\text{ph}} = 3.62$ ms.

Fig. 3 illustrates how we can use the NV center geometry to tune the resonant dipolar-coupling between the spins. Because of the diamond symmetry, we can distinguish 4 groups ("classes") of identical NV centers, based on the four possible orientation of the NV axis in the crystal. These four classes have *a priori* different spin resonance frequency based on the projection of the magnetic field on their respective NV axis. By tuning the angle of the magnetic field, we can therefore control the frequency detuning between the different classes, and as a result we can control the number of resonant NV centers by bringing in and out of resonance several classes.

Fig. 3 (c) in particular shows the dipole-induced lifetime T_1^{dd} for a given class as a function of the energy detuning with another class. One can notice that the

width of the T_1^{dd} bell shape is significantly larger than that of the frequency overlap between the two classes, modeled here by an ODMR line stretched by a factor $\sqrt{2}$ (see SI). This extra width can not be attributed to the dipole-dipole coupling strength : for a 3 ppm concentration of NV^- centers, the average coupling strength between neighboring NV centers should be ~ 27 kHz, two order of magnitude below the observed width. We can also see that T_1^{dd} follows a Lorentzian shape despite the ODMR lines being non-Lorentzian. These two observations corroborate the idea developed in [1] that the interaction width is dominated by the noise spectrum of the lifetime-limited fluctuators.

The depolarization in low field can be explained in large part due to the lift of degeneracy between the four classes, as can be seen in Fig. 4 : On the top part, a magnetic field is scanned in a direction chosen in order to quickly split the transition frequencies of the four NV classes. We can observe an increase of about 4% in the photoluminescence and an increase by a factor of ~ 16 in the spin dipolar-induced lifetime T_1^{dd} as the magnetic field is increased.

However the splitting of the four classes can not explain entirely the spin depolarization in zero field. The bottom part of Fig. 4 shows a similar experiment when the magnetic field is aligned with the [100] crystalline axis. In this particular orientation, the projection of the magnetic field on the four NV classes is identical, meaning that the resonance condition between the NV centers remains the same in zero and non-zero field. Yet we can still observe an increase of $\sim 0.5\%$ in the photoluminescence and an increase of $\sim 30\%$ in T_1^{dd} . We attribute the remaining depolarization to two phenomena : the change of basis in the spin Hamiltonian's eigenstate in zero-field, and the near-resonance of the dipolar double-flip processes.

The decrease of spin lifetime and the associated drop in PL for $B \approx 20$ G is due to cross-relaxations between NV⁻ centers with a ^{13}C as first a neighbor, and those without [4].

The dipole-dipole interaction Hamiltonian between two spins \vec{S}_1 and \vec{S}_2 reads :

$$\mathcal{H}^{\text{dd}} = -\frac{J_0}{r^3} \left(3 \left(\vec{S}_1 \cdot \vec{u} \right) \left(\vec{S}_2 \cdot \vec{u} \right) - \vec{S}_1 \cdot \vec{S}_2 \right), \quad (2)$$

Where $J_0 = (2\pi)52 \text{ MHz} \cdot \text{nm}^3$, \vec{r} is the relative positions of the two spins and $\vec{u} = \frac{\vec{r}}{\|\vec{r}\|}$. In situations such as the one described in Fig. 3, the only relevant terms of \mathcal{H}^{dd} in term of population transfer are the flip-flop terms such as $\langle 0, +1 | \mathcal{H}^{\text{dd}} | +1, 0 \rangle$. In zero magnetic field however, we have to take into account other aspects of the dipolar Hamiltonian in consideration.

First is the change of basis of the single spin Hamiltonian \mathcal{H}_s : in zero external magnetic field, the eigenstates of \mathcal{H}_s in the $\{|+1\rangle, |-1\rangle\}$ manifold are determined by local electric and magnetic field coming from nearby impurities [3]. For the samples used in this study, \mathcal{H}_s was dominated by local electric field (see SI), meaning that

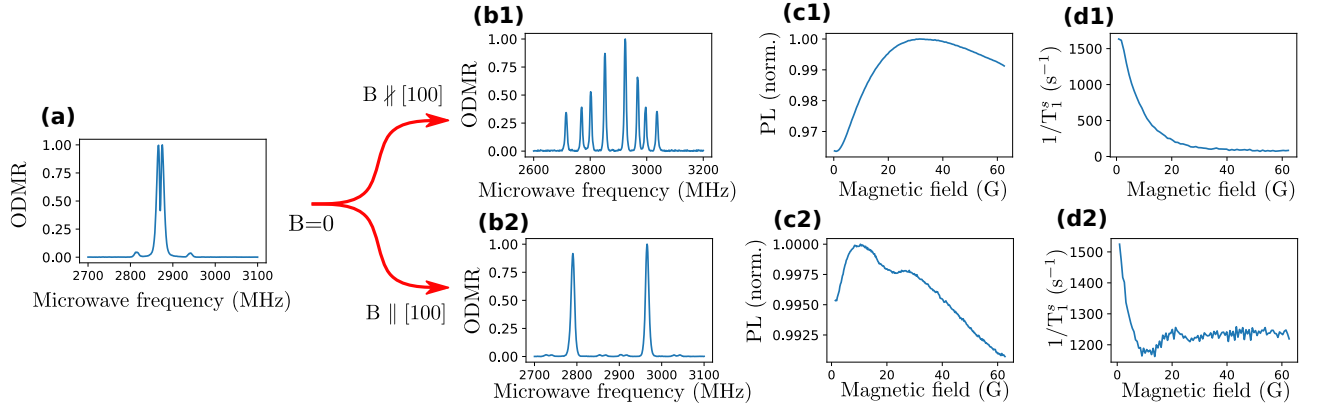


FIG. 4. Dependency of the magnetic field angle for the zero field depolarization. (a) ODMR spectrum in zero field. (b) ODMR spectrum for a magnetic field ≈ 60 G. The field is aligned (b2) or misaligned (b1, $\approx 24^\circ$ of misalignment) with the crystalline [100] direction. (c) Normalized photoluminescence of the NV^- ensemble as a function of the magnetic field amplitude for the same field orientations as fig (b). (d) Stretch part of the lifetime decay as a function of the magnetic field amplitude for the same field orientations as fig (b).

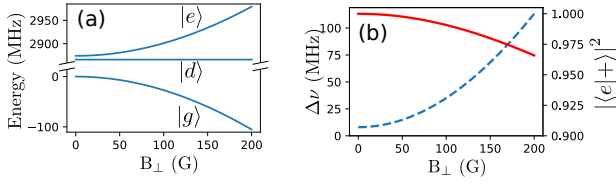


FIG. 5. Simulated eigenstates of the spin Hamiltonian in the presence of purely transverse magnetic field. (a) Energies of the three eigenstates $|g\rangle$, $|d\rangle$ and $|e\rangle$ as a function of the magnetic field amplitude. (b) Blue dashed curve : frequency detuning between the two transitions $|g\rangle \leftrightarrow |d\rangle$ and $|g\rangle \leftrightarrow |e\rangle$. Plain red curve : projection of $|e\rangle$ on $|+\rangle = (|+1\rangle + |-1\rangle)/\sqrt{2}$ as a function of the magnetic field. Is equal to the projection of $|g\rangle$ on $|0\rangle$.

the proper eigenstates of \mathcal{H}_s in zero magnetic field are $\{|0\rangle, |+\rangle = \frac{|+1\rangle + |-1\rangle}{\sqrt{2}}, |-\rangle = \frac{|+1\rangle - |-1\rangle}{\sqrt{2}}\}$

Because the splitting between $|+\rangle$ and $|-\rangle$ (≈ 9 MHz) is much greater than the dipole-dipole interaction ($J_0/r^3 \approx 30$ kHz), the flip-flop terms of \mathcal{H}^{dd} now reads $\langle 0, + | \mathcal{H}^{\text{dd}} | +, 0 \rangle$. Averaging over all relative positions of two aligned NV centers, we found that on average $|\langle 0, + | \mathcal{H}^{\text{dd}} | +, 0 \rangle|$ was greater than $|\langle 0, + | \mathcal{H}^{\text{dd}} | +, 0 \rangle|$ by a factor $1.8 \sim 2$ depending on the correlation lengths of the local electric field. When all four classes are resonant, the theoretical decrease in the spin lifetime due to the change of basis is $\sim 20\%$ (See SI de ouf).

Second is the near-resonance condition of the double-flip processes such as $\langle 0, + | \mathcal{H}^{\text{dd}} | -1, 0 \rangle$ or $\langle 0, + | \mathcal{H}^{\text{dd}} | -, 0 \rangle$. These terms usually couple non resonant states due to the energy mismatch between $|+1\rangle$ and $|-1\rangle$. However in zero field, the energy splitting between $|+\rangle$ and $|-\rangle$ is comparable with the fluctuator's spectral width measured in Fig. 3 to be ≈ 8 MHz, making the double-flip population transfer possible.

In order to evaluate the relative contribution of these

two factors, we investigate the spin relaxation in the case of pure transverse magnetic field. Calling $\{|g\rangle, |d\rangle, |e\rangle\}$ the three eigenstates of the spin Hamiltonian in the presence of purely transverse magnetic field, Fig. 5 shows that for small enough magnetic field, the $\{|g\rangle, |d\rangle, |e\rangle\}$ basis is almost equal to the $\{|0\rangle, |+\rangle, |-\rangle\}$ basis : for $B = 120$ G, $\langle 0 | g \rangle \approx 0.99$, $\langle -1 | d \rangle = 1$ and $\langle +1 | e \rangle \approx 0.99$. However the energy difference between $|d\rangle$ and $|e\rangle$ reaches ≈ 45 G which is high enough to cancel out the double-flip processes and allows us to isolate the change of basis hypothesis from the double-flip one.

Fig. 6 shows the dipole-dipole spin relaxation T_1^{dd} for a class of NV whose axis is orthogonal to the applied magnetic field. By monitoring the frequencies of the transitions $|g\rangle \leftrightarrow |d\rangle$ and $|g\rangle \leftrightarrow |e\rangle$, we can deduce the detuning $\Delta\nu$ between the two-spin states $|0, +\rangle$ and $|-, 0\rangle$. Plotting then $1/T_1^{\text{dd}}$ as a function of $\Delta\nu$, we can notice two things.

First the decrease of $1/T_1^{\text{dd}}$ as $\Delta\nu$ increases corresponding to the loss of effectiveness of the double flip processes. The shape of the curve matches decently well with the tail of a Lorentzian centered in $\Delta\nu = 0$ with a width of 8 MHz, confirming that the double-flip process interaction range is likely determined by the fluctuators noise spectrum just like the flip-flop processes.

Second is the plateau reached for $B \gtrsim 30$ MHz. The final $1/T_1^{\text{dd}}$ value is about twice as low as that for a spin well aligned with the magnetic field, showing the increased depolarization due to the change of basis.

This experiment shows that, for the sample studied here, the depolarization due to the double-flip processes dominates the one due to the change of basis in zero magnetic field.

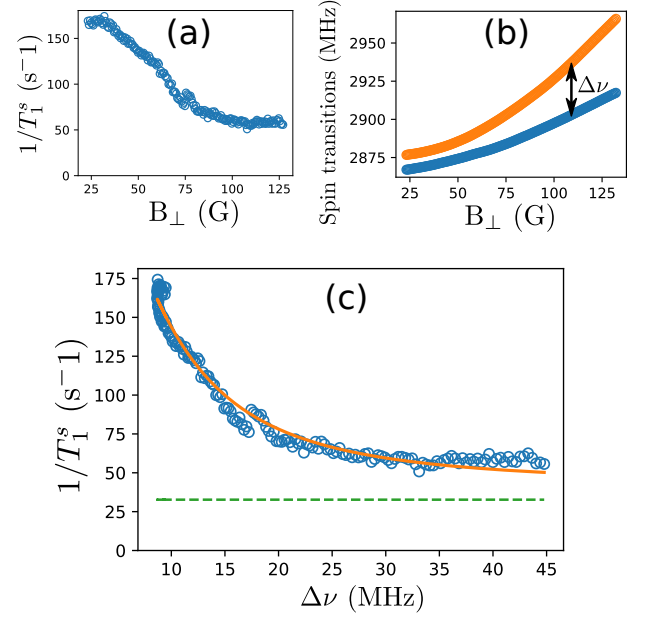


FIG. 6. Modification of the stretch lifetime in the presence of purely transverse magnetic field. (a) Stretch component of the ensemble lifetime as a function of the field amplitude. (b) Measured transition frequencies through ODMR spectrum. (c) Stretch component of the lifetime as a function of the frequency detuning between the two transitions (blue circles), fitted by a Lorentzian centered in $\Delta\nu = 0$ with half width at half maximum 8MHz. The green dashed line correspond to the lifetime of a single class aligned with the magnetic field.

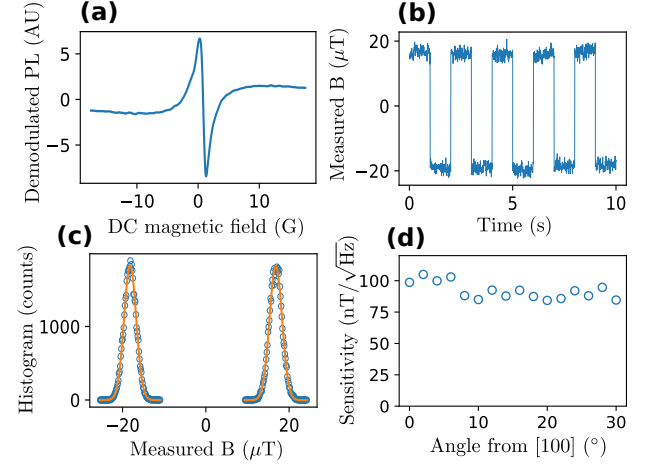


FIG. 7. Low field magnetometry protocol. (a) Demodulated photoluminescence as a function of an eternally applied magnetic field with an additional oscillatory magnetic field. (b) Measured magnetic field when alternating a small external magnetic field offset. (c) Histogram of the measurement in Fig. (b) fitted with gaussians of standard deviation $\sigma = 1.5 \mu\text{T}$

MAGNETOMETRY

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