

SUPPLEMENTARY MATERIAL

Depolarization of interacting NV^- centers at low magnetic fields

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I. NV^- GROUND STATE HAMILTONIAN UNDER LOW MAGNETIC FIELD

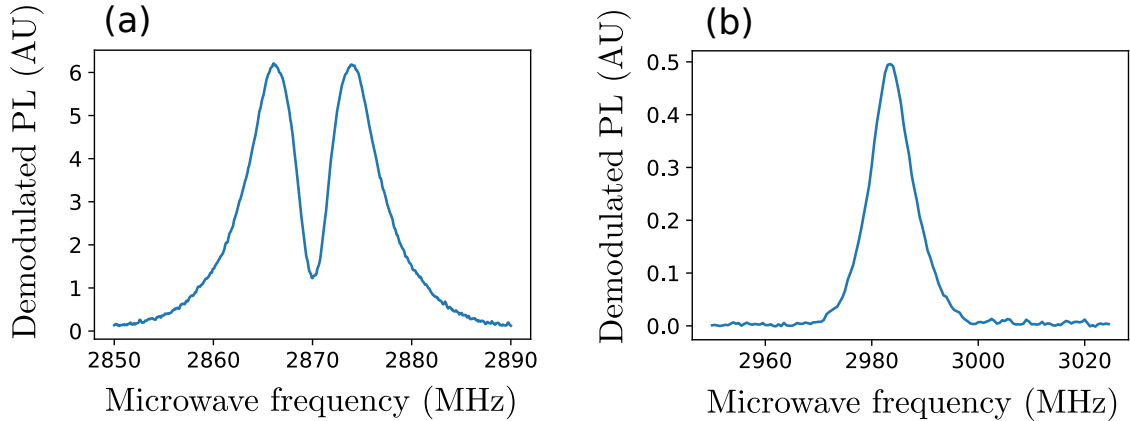


FIG. 1. ODMR measurement (a) under zero magnetic field, (b) under a magnetic field of 50 G when zooming on a single NV class.

Under zero external magnetic field, there are three possible processes that can cause a splitting of the $\{|+1\rangle, |-1\rangle\}$ states : local electric fields, crystal strain and local magnetic fields. Of these three causes, only the electric field can explain the shape of the ODMR line that is observed under zero external magnetic field (Fig. 1a) [8].

Indeed, random local magnetic field would produce a single broadened line while crystal strain would shift the zero field splitting (ZFS) by the same order of magnitude as the splitting between the levels, which would merge the two transitions onto a single line. Due to the large difference between the longitudinal and transverse electric field susceptibilities ($d_{\parallel} = 0.35 \text{ Hz cm/V}$ and $d_{\perp} = 17 \text{ Hz cm/V}$ [11]), random local electric field can, on average, cause a splitting that is much stronger than the ZFS shift and result in a two peak spectrum centered on 2.87 GHz.

In our model for dipole-dipole coupling under small magnetic fields, we will therefore neglect the contribution of the strain, local magnetic fields and longitudinal electric fields.

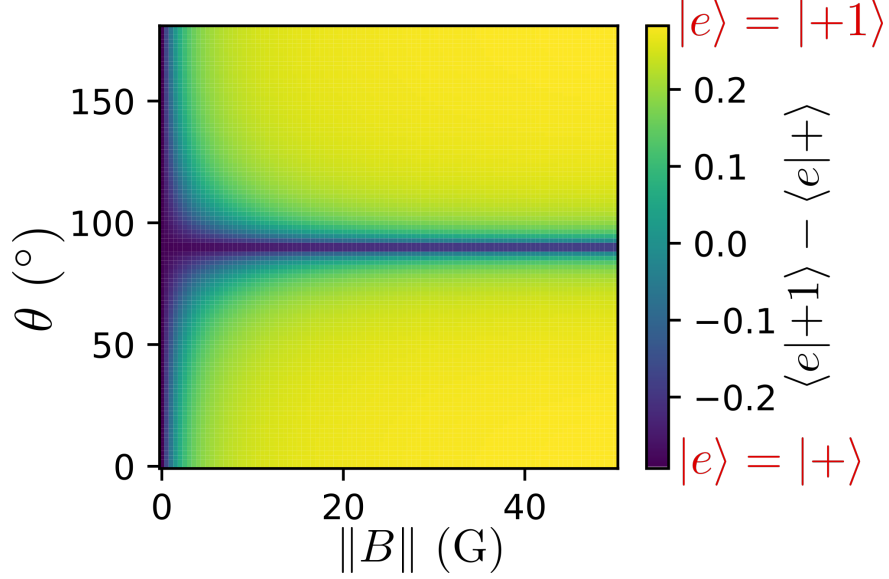


FIG. 2. Numerical simulations showing the closeness of the Hamiltonian second excited state $|e\rangle$ with the states $|+1\rangle$ and $|+\rangle$, as a function of the magnetic field amplitude and angle θ with respect to the NV axis. A value of $d_{\perp}E_{\perp} = 4$ MHz was chosen.

We also do not take into account the hyper-fine structure of the NV center because of the large inhomogeneous broadening of the transitions in all our samples (Fig. 1b). We will then consider the following spin Hamiltonian for the NV^- ground state :

$$\mathcal{H}_s/h = DS_z^2 + \gamma_e \mathbf{B}_{\text{ext}} \cdot \mathbf{S} + d_{\perp} [E_x(S_y^2 - S_x^2) + E_y(S_x S_y + S_y S_x)], \quad (1)$$

where $D = 2.87$ GHz is the zero field splitting and $\gamma_e = 2.8$ MHz/G the gyromagnetic ratio of the electron.

In the absence of an external magnetic field, the symmetry of the Hamiltonian in the (xy) plane allows us to pick the x direction along the electric field. The eigenstates of \mathcal{H}_s then become $\{|0\rangle, |+\rangle = \frac{|+1\rangle + |-1\rangle}{\sqrt{2}}, |-\rangle = \frac{|+1\rangle - |-1\rangle}{\sqrt{2}}\}$.

In the presence of a magnetic field at an angle θ with respect to the NV axis, we denote the Hamiltonian eigenstates $\{|g\rangle, |d\rangle, |e\rangle\}$ in ascending order of energy. Fig. 2 shows how close the $|e\rangle$ state is to the $|+1\rangle$ and $|+\rangle$ states as a function of the external magnetic field and angle θ . A similar analysis for the $|d\rangle$, $|-1\rangle$ and $|-\rangle$ states show similar results, while $|g\rangle$ is pretty much equal to $|0\rangle$ for $B < 100$ G.

This result tells us that, in most cases, the $\{|0\rangle, |+\rangle, |-\rangle\}$ basis is only the good Hamiltonian basis for magnetic field smaller than a few Gauss, except in the case of pure transverse

magnetic field where the $\{|0\rangle, |+\rangle, |-\rangle\}$ basis remains a good basis even for sizable magnetic fields.

II. SAMPLES

Here are the various samples used in this study :

- **HPHT-150-1** : A 150 μm HPHT 1b diamond irradiated and annealed to reach a concentration $[\text{NV}^-] \approx 3$ ppm. This sample was bought from Adamas Nanotechnology (MDNV150umHi). It is used in Fig. 2 (b), 2(d) and 3 of the main text, as well as Fig. 1, 4 and 5 of the SI.
- **HPHT-150-2** : Another sample from the same batch as HPHT-150-1. This sample is used in Fig. 4 of the main text and Fig. 6 (c) of the SI.
- **HPHT-150-3** : Another sample from the same batch as HPHT-150-1. This sample is used in Fig. 9 of the SI.
- **HPHT-15-1** : A 15 μm diamond with similar properties as the last ones, also bought from Adamas Nanotechnology (MDNV15umHi). This sample is used in Fig. 5 of the main text.
- **HPHT-15-2** : Another sample from the same batch as HPHT-15-1. This sample is used in Fig. 12 and 13 of the SI.
- **HPHT-1-1** : A 1 μm diamond with similar properties as the last ones, also bought from Adamas Nanotechnology (MDNV1umHi). This sample is used to perform a sensitivity measurement similar to the one done in Fig. 5 of the main text.
- **CVD-1** : A CVD type IIa bulk diamond. This sample has not been irradiated and contains $[\text{NV}^-] \approx 50$ ppb. This sample is used in Fig. 2(a) of the main text.
- **CVD-2** : A CVD bulk diamond described in [10]. This one has been irradiated and annealed and contains $[\text{NV}^-] \approx 4$ ppm. This sample is used in Fig. 10 of the SI.

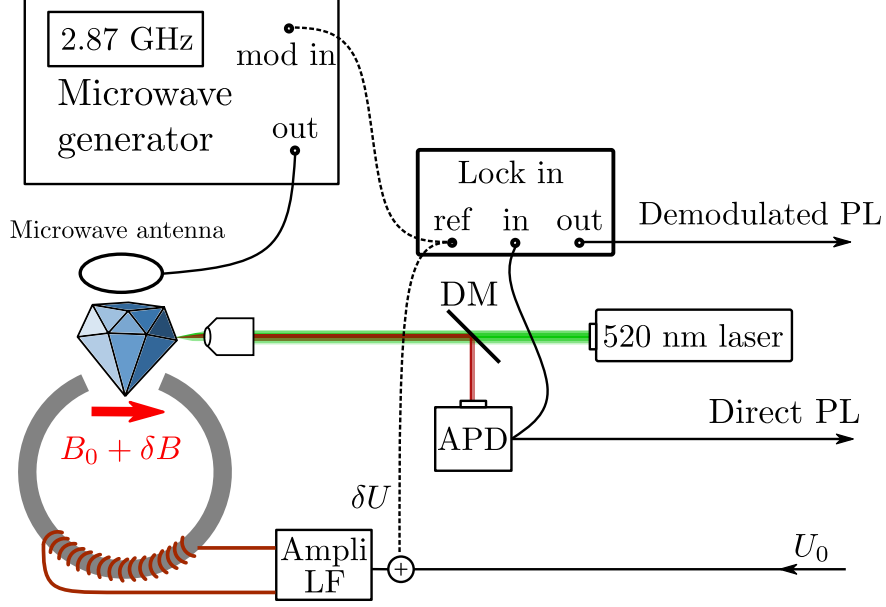


FIG. 3. Experimental Setup.

III. EXPERIMENTAL SETUP

Fig. 3 shows the general purpose experimental setup used for all the experiments presented in this article.

The optical polarization and readout of the spins is done by focusing a green laser on the diamond sample with an objective lens ($\text{NA}=0.65$), and collecting the back-scattered red fluorescence from the NV center on an avalanche photo-diode (APD, Thorlabs APD410A). The laser is filtered out using a dichroic mirror and a notch filter. The NV^0 fluorescence is filtered using an additional 645 nm longpass filter.

The laser used here is a Picoquant PDL 800-D with a 520 nm LDH laser head, providing pulses of 40 ps at a rate of 20 MHz, with an average power of $0.5 \sim 5$ mW. These fast pulses serve no purpose in the actual experiment. This laser was used solely because of its high amplitude stability. The trigger of the pulses is generated externally in order to achieve fast gating of the laser for the T_1 measurement. We previously did similar experiment using a continuous 532 nm laser and observed no difference in the spin response.

The magnetic field is provided by a homemade electromagnet composed of a c-shape iron core and copper wires. The magnet is mounted on two mechanical rotation stages, allowing a control on the polar and azimuthal angle of the magnetic field within a fraction of a degree and is alimented through a low frequency amplifier (Leybold power function generator 522

63).

The microwave field is generated by a Rhode & Shwarz SMB 100A and is emitted with a handmade loop antenna. The microwave field is gated by a Mini-circuits ZASWA-2-50DRA+ switch controlled externally and amplified by a Mini-circuits ZHL-5W-422+ amplifier.

A lock-in amplifier (SRS SR830 DSP) is used either to modulate the microwave amplitude for ODMR measurement, or to add an oscillatory magnetic field for the magnetometry protocol. In both case we use a modulation frequency ~ 1 kHz and demodulate the APD signal.

We did not use magnetic shielding to protect from the earth magnetic field since the splitting due to B_{earth} is lower than the splitting due to the local electric field for the samples used here : $\gamma_e B_{\text{earth}} \approx 1.5 \text{ MHz} < d_{\perp} E_{\perp} \approx 4 \text{ MHz}$.

IV. EXPERIMENTAL DETAILS

A. T_1 fitting Protocol

Fig. 2-c) of the main text shows the sequence employed for measuring T_1^{dd} . 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 . In highly doped samples, this sequence often results in artifacts, mostly due to charge state transfer in the dark [2, 4, 5]. It is therefore convenient to repeat the sequence with an additional π pulse on one of the eight NV spin-resonances right before the spin read-out to prepare the remaining $|0\rangle$ polarization into a darker $|+1\rangle$ or $|-1\rangle$ state [2, 7, 9]. 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.

Fig. 4 shows the result of a lifetime measurement for the two scenarios presented in main text, namely $B = 0$ and $B = 50$ G in a direction for which every classes are split, on sample HPHT-150-1. In main text, we fitted these results with the expression

$$S(\tau) = A \exp\left(-\sqrt{\frac{\tau}{T_1^{\text{dd}}}} - \frac{\tau}{T_1^{\text{ph}}}\right), \quad (2)$$

in order to extract the T_1 part associated with the dipole-dipole coupling, as was previously done to study the cross-relaxation between NV and substitutional nitrogen (P1) center [6].

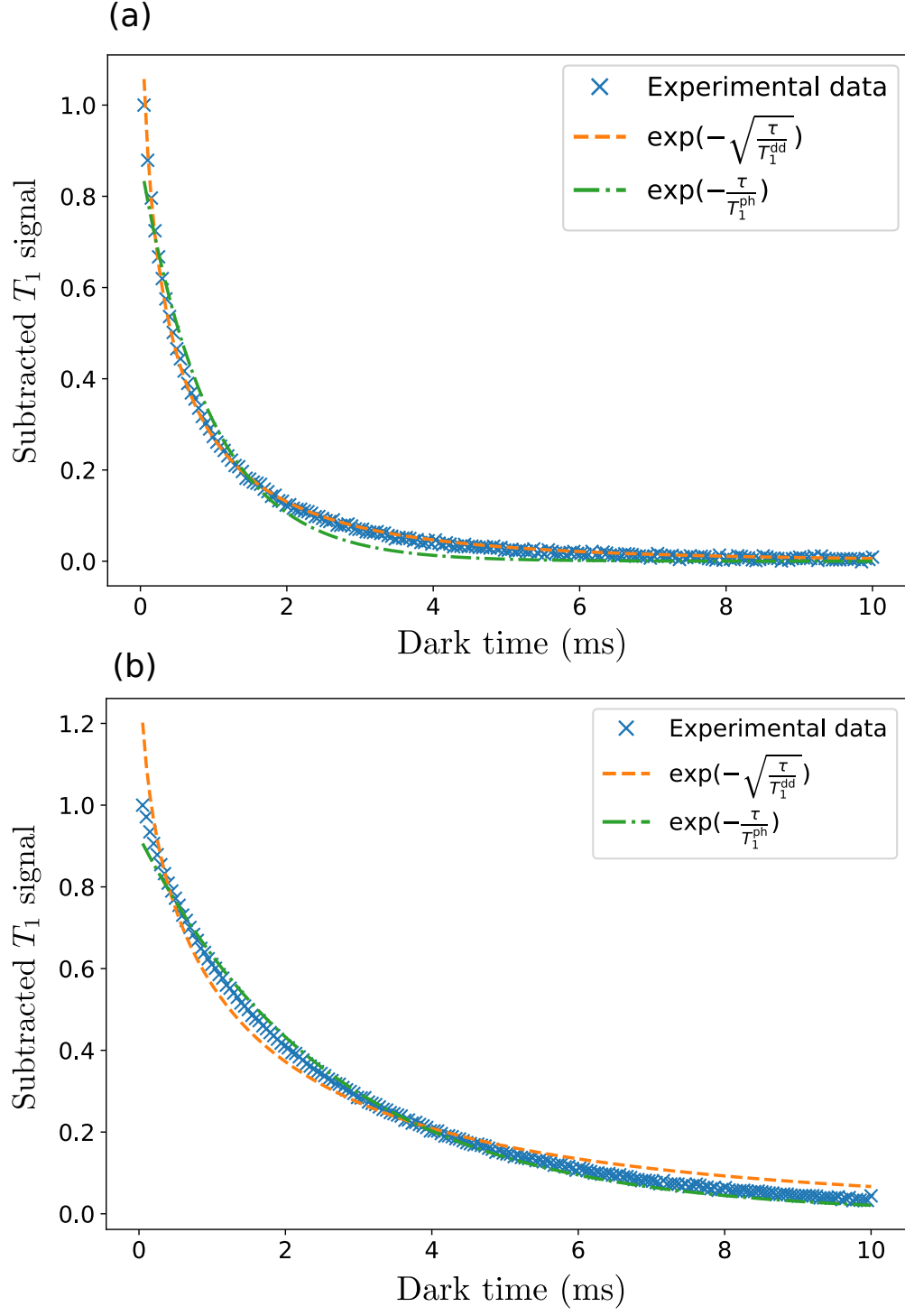


FIG. 4. T_1 measurement with purely exponential and purely stretched exponential fits (a) in zero magnetic field (b) in non-zero magnetic field.

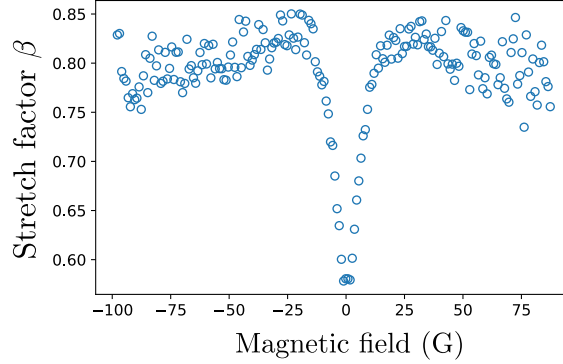


FIG. 5. Best stretch factor β for a T_1 fit of the form $f(\tau) = A \exp\left(-\left(\frac{\tau}{T_1}\right)^\beta\right)$ as a function of an arbitrarily oriented magnetic field amplitude.

In Fig. 4, we show what happens when we fit the T_1 measurements with either purely exponential or purely stretched exponential fits. We can see that for $B = 0$, which corresponds to the regime where dipole-dipole relaxation is strong, the experimental data is closely fitted by the stretched exponential profile, whereas for $B = 50$ G, the experimental data follows more closely the exponential profile. In order to describe both regime simultaneously, we have to take both T_1^{dd} and T_1^{ph} into account in our analysis.

Since we are only interested in the stretched exponential part of the lifetime decay, and we are assuming that the phonon-limited exponential decay does not depend on the external magnetic field, we decided to fix the value T_1^{ph} for each sample, in order to reduce the number of free parameters in eq. 2. The value chosen for all T_1 measurements on samples HPHT-150-1 and HPHT-150-2 was $T_1^{\text{ph}} = 3.62$ ms.

Another possibility to fit our T_1 measurements which yields satisfactory fits is to use an arbitrary stretched factor in the fitting function :

$$S(\tau) = A \exp\left(-\left(\frac{\tau}{T_1}\right)^\beta\right). \quad (3)$$

Fig. 5 shows the optimal β parameter as a function of the external magnetic field on sample HPHT-150-1, which confirms that the T_1 profile gets closer to a purely stretched exponential in zero magnetic field.

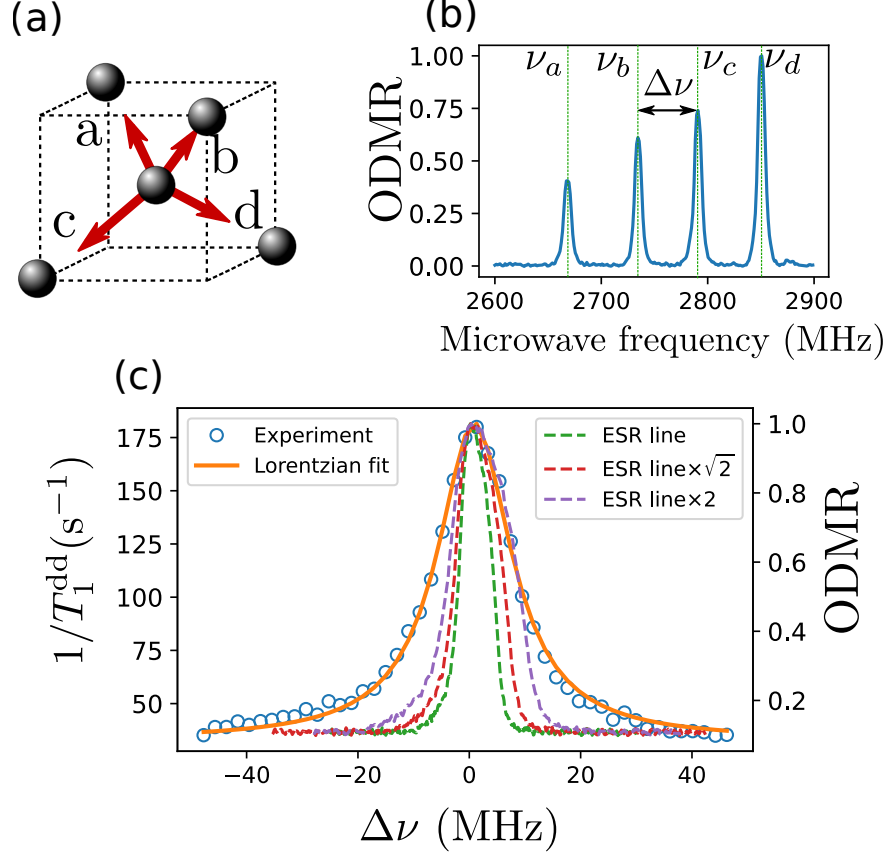


FIG. 6. Dipole-dipole depolarization for two near-resonant classes. (a) Sketch of the four possible NV orientations (“classes”) in a single diamond. (b) ODMR spectrum showing four $|0\rangle \rightarrow |-1\rangle$ resonances corresponding to the four spin classes. The detuning $\Delta\nu$ between the classes b and c was controlled by changing the orientation of the external magnetic field. (c) Stretched part of the lifetime decay curve for the spins resonant with ν_c as a function of the detuning $\Delta\nu$ (blue circles), fitted by a Lorentzian with half width at half maximum 8.04 MHz. Single class ESR line stretched by a factor of 1, $\sqrt{2}$ and 2 are added for comparison.

B. Spectral range of the dipole-dipole cross-relaxations

An experimental signature of the fluctuator hypothesis developed in [2] is the dependence of the depolarization rate when two near-resonant classes are brought to resonance : if there are indeed very fast decaying NV centers (fluctuators with lifetime $T_1^f < 100$ ns), then the spectral width of the fluctuator would be greater than $1/T_2^*$. For simplicity, it is typically assumed that the latter is the same for all spins in the crystal. This means that NV centers would be able to exchange spin quanta (flip-flop) with non-resonant NV fluctuators detuned

by $\Delta\nu$ such that $2\pi/T_1^f > \Delta\nu > 2\pi/T_2^*$.

In order to verify this, we measure the spectral overlap between two classes, which in the absence of fluctuator should be on the order of $1/T_2^*$, and compare it to the actual width of the depolarization rate of the spins as a function of detuning.

Fig. 6 shows the results of such an experiment. We measured the stretched part of the NV's decay rate at each detuning and obtained Fig. 6 -c), which is very well fitted by a Lorentzian of half-width $\sigma = 8.04$ MHz. We should note that this broadening can not be explained by the direct coherent dipole-dipole interaction induced splitting : for a sample with 3 ppm of NV centers, the average dipole-dipole interaction strength between two nearest NV centers $J_0/r^3 \sim 27$ kHz which is several order of magnitude lower than the broadening we observe.

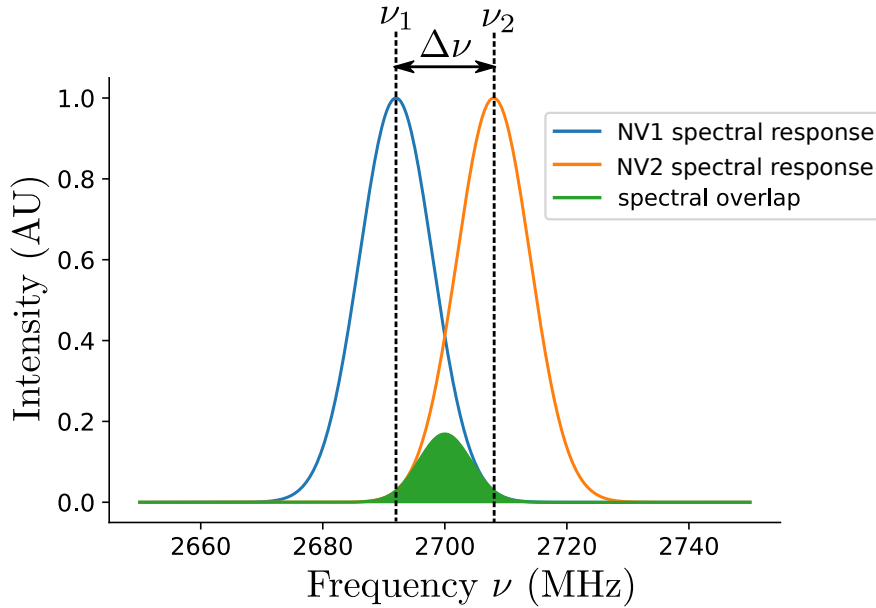


FIG. 7. Illustration of the spectral overlap for two gaussian spectra.

We can compare this obtained width to the width of an ODMR line of a single class of NV centers, stretched by a factor of $\sqrt{2}$ and 2 to simulate the spectral overlap of two classes. One should notice that, not only is $1/T_1^{\text{dd}}$ significantly larger than the ODMR profile, it also does not have the same shape.

As illustrated in Fig. 7, we define the spectral overlap $S(\Delta\nu)$ between two spins of spectral response $S_1(\nu)$ and $S_2(\nu)$, centered respectively on the frequencies ν_1 and ν_2 where

$\Delta\nu = \nu_2 - \nu_1$ as :

$$S(\Delta\nu) = \int S_1(\nu, \nu_1) S_2(\nu, \nu_2) d\nu. \quad (4)$$

In order to approximate the spectral overlap in our experiment, we will consider the analytical solution in the Gaussian and Lorentzian case :

- For two gaussians of standard deviation σ , the spectral overlap as a function of the detuning $\Delta\nu = \nu_1 - \nu_2$ is itself a gaussian of standard deviation $\sigma' = \sqrt{2}\sigma$. :

$$\begin{aligned} S(\Delta\nu) &\propto \int \exp\left(-\frac{(\nu - \nu_1)^2}{2\sigma^2}\right) \exp\left(-\frac{(\nu - \nu_2)^2}{2\sigma^2}\right) d\nu \\ &\propto \exp\left(-\frac{(\Delta\nu)^2}{4\sigma^2}\right). \end{aligned}$$

- For two Lorentzian profile with width σ , the overlap function is itself a Lorentzian with width $\sigma' = 2\sigma$:

$$\begin{aligned} S(\Delta\nu) &\propto \int \frac{1}{1 + \frac{(\nu - \nu_1)^2}{\sigma^2}} \cdot \frac{1}{1 + \frac{(\nu - \nu_2)^2}{\sigma^2}} d\nu \\ &\propto \frac{1}{1 + \frac{(\Delta\nu)^2}{4\sigma^2}}. \end{aligned}$$

The ODMR lines that we measure are neither Lorentzian nor Gaussian (although they tend to be closer to Gaussians), and can even be asymmetric. Nevertheless, the overlap between two classes can most likely be approximated by a single class ODMR profile stretched by a factor between $\sqrt{2}$ and 2.

C. Lift of the degeneracy between the four classes

We give here an estimate of the magnetic field required to lift the degeneracy of the four classes in the case of Fig. 3(b) of the main text. Fig. 8 shows the simulated energies for the 4 $|0\rangle \rightarrow |-1\rangle$ and the 4 $|0\rangle \rightarrow |+1\rangle$ transitions as a function of the magnetic field, and the difference in energy between the closest pairs of transitions. We can see that the energy detuning between the classes $\Delta\nu$ crosses the value of the dipole-dipole interaction range (8.04 MHz, measured on sample HPHT-150-2 but likely to be similar on sample HPHT-150-1) for magnetic field values $\|B\| = 12 \sim 15$ G, which are close to the half-width of the zero-field feature in Fig. 3(b)(ii) and (iii) of the main text.

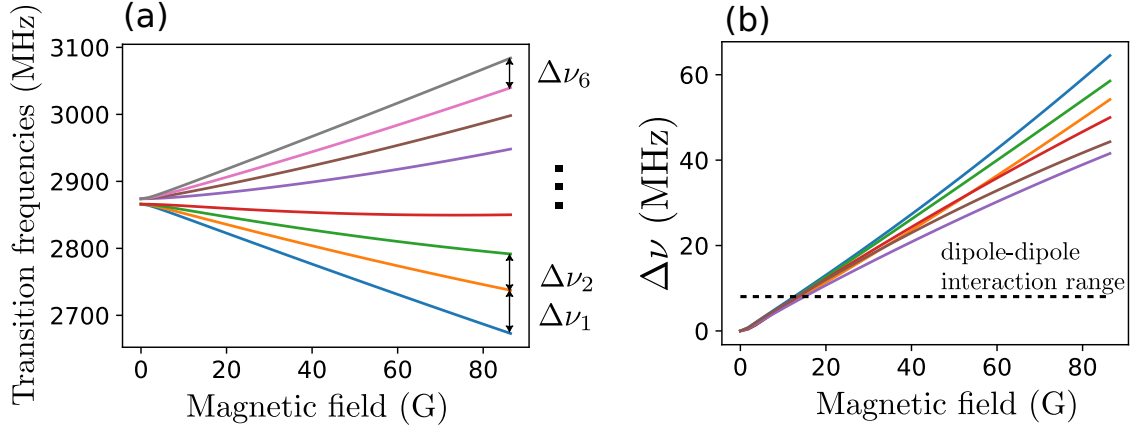


FIG. 8. (a) : Simulation of the 8 possible transition frequencies (four $|0\rangle \rightarrow |-1\rangle$ and four $|0\rangle \rightarrow |+1\rangle$ transitions) as a function of the magnetic field amplitude for the orientation of the magnetic field shown in Fig. 3(b) of the main text. (b) Difference in frequency between each pair of closest classes as a function of the magnetic field amplitude. The dotted line $y=8.04$ MHz corresponds to an estimation of the dipole-dipole interaction frequency range.

D. Effect of laser polarization

Previous studies [1, 3] reported the presence of a photoluminescence dip in zero magnetic field which was heavily dependent on the laser polarization angle with respect to the diamond axes and the magnetic field.

We did not observe a strong dependence on the laser polarization with the samples used in this study. Fig.9 shows the photoluminescence from one of our sample as a function of the magnetic field, either directly or with a modulation of the magnetic field, for 6 polarization angles and saw no major difference.

In particular, we did not observe the apparition of an anti-line inside the main dip, unlike what was observed in the two previously cited work. We expect that the main reason behind the different behaviors is that we seem to observe far greater spin depolarization in zero-field, which we attribute to dipole-dipole coupling. This effect could hide smaller effects such as the one related to the laser polarization.

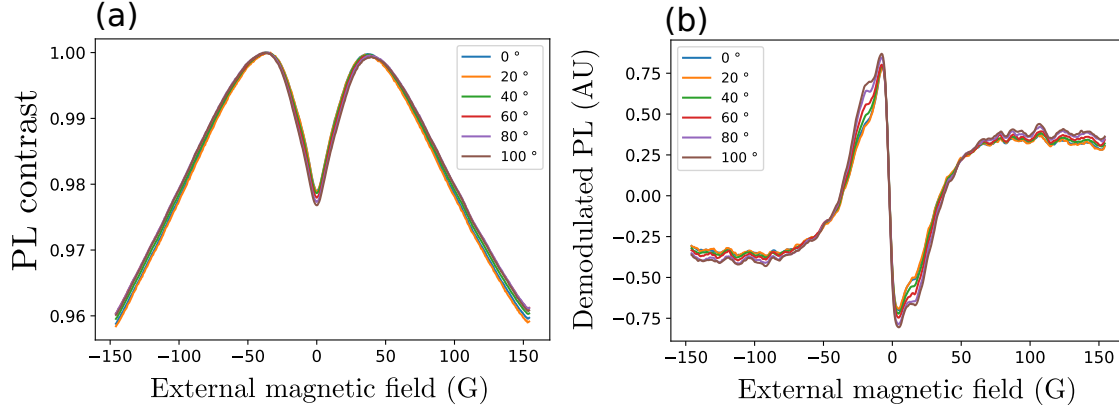


FIG. 9. Effect of the polarization of the incident laser. (a) Photoluminescence of sample HPHT-150-3 as a function of arbitrarily oriented magnetic field amplitude for various polarization angle. (b) Demodulated PL in the same conditions.

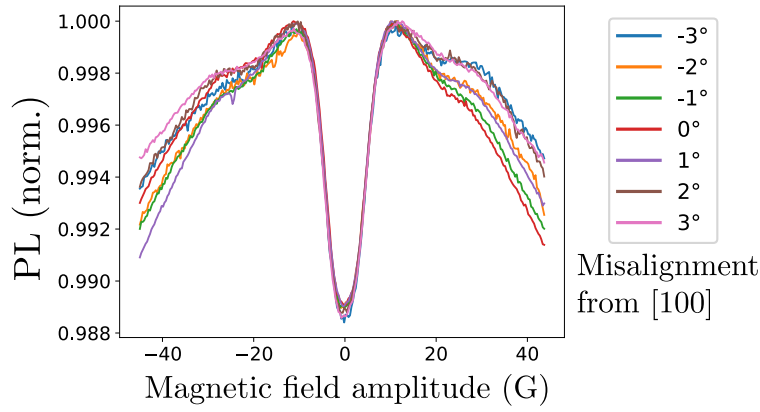


FIG. 10. PL of sample CVD-PPM as a function of the external magnetic field amplitude for different magnetic field misalignments with respect to the $[100]$ axis.

E. Alignment of the magnetic field along $[100]$ axis

We claim that the decrease in PL at low magnetic field in Fig. 3 (c)(ii) of the main text is due to the local electric field and the double flip processes. One other possibility would be a misalignment of the magnetic field with respect to the $[100]$ axis as it is scanned. The four classes could for instance be truly resonant only with zero external magnetic field and not when the B field is large.

Fig. 10 shows the change in PL with respect to the external magnetic field amplitude for various misalignments of the magnetic field direction compared to the $[100]$ diamond

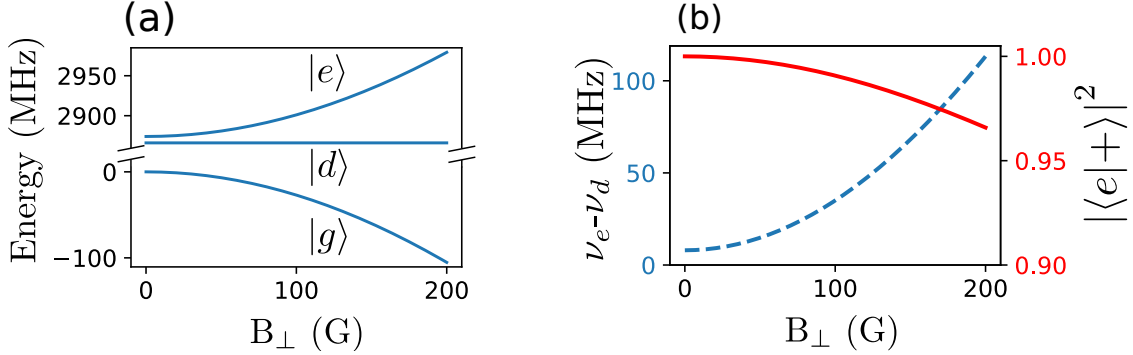


FIG. 11. (a) Simulated energy level of the three eigenstates $|g\rangle$, $|d\rangle$ and $|e\rangle$ for an NV center under purely transverse magnetic field. The spin Hamiltonian is given in eq. 1 and a value of $d_{\perp}E_{\perp} = 4$ MHz was chosen (b) Blue dashed curve : frequency splitting between the $|d\rangle$ and $|e\rangle$ states as a function of the transverse magnetic field, red plain line : matching factor $|\langle e|+\rangle|^2$ between the $|e\rangle$ and $|+\rangle$ states.

crystalline axis of the sample CVD-2. We can see that the central drop in PL is not critically affected by the misalignment of the magnetic field, and we are therefore confident that the drop in PL we observe when $B \parallel [100]$ is not an effect of misalignment.

The initial alignment of the magnetic field was controlled by applying a field of ~ 50 G and monitoring the overlap of the four classes with ODMR spectra (maximizing the PL drop is also an effective method). We estimate the initial alignment precision to be $\pm 1^\circ$. The misalignment was then introduced by rotating the electromagnet along an arbitrary axis. The slight asymmetry between the positive and negative value of the magnetic field could come from the earth magnetic field or from hysteresis in the magnetic core of our electromagnet.

F. Emulation of the electric field with transverse magnetic field

In the main text we use the property that, under purely transverse magnetic field, the eigenstates of the single spin Hamiltonian are close to the $\{|0\rangle, |+\rangle, |-\rangle\}$ basis, described in the main text and in sec. I of these supplementary. We will discuss here the validity of this claim.

We will call the true eigenstates of the single spin Hamiltonian under purely transverse magnetic field $\{|g\rangle, |d\rangle, |e\rangle\}$ in ascending order of energy. The $|d\rangle$ state is strictly equal to

the $|-\rangle$ state. The $|g\rangle$ and $|e\rangle$ states can be written as a mixture of the $|0\rangle$ and $|+\rangle$ states, and their exact expression depends on the field value. As B_\perp approaches 0 however, $|g\rangle$ and $|e\rangle$ tend toward $|0\rangle$ and $|+\rangle$ respectively.

Fig. 11 (a) shows a simulation of the energy levels of the three eigenstates $|g\rangle$, $|d\rangle$ and $|e\rangle$ as a function of the transverse magnetic field amplitude. Of importance for the results in main text is the splitting $\Delta\nu$ between the $|e\rangle$ and $|d\rangle$ states, which is equal to the difference in the transition frequencies $|g\rangle \rightarrow |d\rangle$ and $|g\rangle \rightarrow |e\rangle$ (reported experimentally as ν_+ and ν_- in main text).

This value $\Delta\nu$ is reported on Fig. 11 (b) along with a matching factor $|\langle e|+\rangle|^2$. This factor is always equal to $|\langle g|0\rangle|^2$ since $\langle d|-\rangle = 1$. We can see that indeed, for $B_\perp < 150$ G, $|\langle e|+\rangle|^2 > 0.98$. Crucially, for $B_\perp = 150$ G, the splitting $\Delta\nu$ is equal to ≈ 70 MHz, almost an order of magnitude greater than the dipole-dipole interaction range computed on sec. IV B. This means that we can effectively quench the double-flips by splitting the $|e\rangle$ and $|d\rangle$ states, while still maintaining eigenstates almost equal to $|0\rangle$, $|+\rangle$ and $|-\rangle$.

V. EXTENSION OF THE FLUCTUATOR MODEL TO LOW MAGNETIC FIELDS

A. Results

Here are the results of the fluctuator model developed in [2] that we extended to include double flip as well as local electric fields. The following sections will detail the calculations that led to these results. Let us first summarize our theoretical findings :

- The dipole-induced spin decay is lowest for a spectrally isolated class of NV centers when the transverse magnetic field is negligible. In this case the dipole-dipole relaxation is limited by flip-flop processes between spins of the same class. We will call the corresponding stretched lifetime $T_1^{\text{dd}} \equiv T_0$.
- For a single class, still spectrally isolated from the three other classes, but this time dominated by the electric field or transverse magnetic field (as long as $\gamma_e B_\perp \ll D$), the change in the eigenstates of the single particle Hamiltonian results in an increase of the average flip-flop rate. This leads to a new theoretical stretched lifetime $T_1^{\text{dd}} = 4 T_0$.

- When a magnetic field is applied along the $[100]$ axis (for $5 \text{ G} < |B| \ll D/\gamma_e$), all four classes are resonant and dominated by the longitudinal part of the magnetic field. This results, in increase in the average flip-flop rate compared to the isolated class case, and the predicted stretched exponential lifetime is $T_1^{\text{dd}} \approx 42.8 T_0$.
- In zero external magnetic field, all four classes are also resonant but we assume that the single particle Hamiltonian of each spin is dominated by a random local electric field. The averaging of flip-flop in this case yields a stretched lifetime $T_1^{\text{dd}} \approx 51.4 T_0$, which is about 20% higher than the predicted lifetime for $B \parallel [100]$.

The effect of the double flip processes was not studied quantitatively since the process is never fully resonant and is therefore highly dependent on the initial splitting of the $|+\rangle$ and $|-\rangle$ states. Nevertheless, the presence of double-flip as an additional relaxation channel can only increase the dipole-induced spin decay rate.

B. Summary of the NV-fluctuator model

Here are the main hypotheses and conclusion of the NV-fluctuator model developed in [2] :

The NV^- centers in the crystal are divided between two categories : “normal” NV centers (simply called NV) which, in the absence of dipole-dipole coupling would have a phonon-limited T_1 ($T_1^{\text{NV}} \sim \text{ms}$), and fluctuators who are NV centers with an additional, fast, depolarization mechanism, such that their lifetime $T_1^f < 100 \text{ ns}$. Having such a short lifetime, the fluctuators are almost unpolarized by the green laser, making them invisible in standard optical NV measurement protocol (T_1 , ODMR, etc.).

Assuming a homogeneous distribution of the fluctuators in the bulk of the crystal, the authors of [2] show that the fluctuators create an additionnal decay channel for the NV population, through dipole-dipole interaction, characterized by a decay rate γ which follows the probability distribution :

$$\rho(\gamma) = \frac{e^{-1/(4\gamma T)}}{\sqrt{4\pi\gamma^3 T}}, \quad (5)$$

where the timescale T is defined as

$$\frac{1}{T} = \left(\frac{4\pi n_f J_0 \bar{\eta}}{3} \right)^2 \frac{\pi}{\gamma_f}, \quad (6)$$

where n_f is the fluctuator density in the crystal in nm^{-3} , $J_0 = 52 \text{ MHz} \cdot \text{nm}^3$ is the characteristic dipole-dipole strength between two spins, γ_f is the fluctuator intrinsic decay rate and $\bar{\eta}$ is a dimension-less number which characterizes the average dipole-dipole interaction between the NV centers and the fluctuators (resonance conditions, relative orientations etc. Further details are given below).

The polarization dynamics of the averaged ensemble of NV centers then follows :

$$P(t) = \int_0^\infty \rho(\gamma) e^{-\gamma t} d\gamma = e^{-\sqrt{t/T}}. \quad (7)$$

Which corresponds to the stretched-exponential part of the lifetime measurement.

C. Dipole-Dipole Hamiltonian between two NV^- centers

Let us consider the hamiltonian of an NV center interacting with a fluctuator :

$$\mathcal{H}_{\text{tot}} = \mathcal{H}_1 + \mathcal{H}_2 + \mathcal{H}_{\text{dd}}. \quad (8)$$

\mathcal{H}_1 and \mathcal{H}_2 are the single particle Hamiltonian of the NV and fluctuator, described by the equation (1), and \mathcal{H}_{dd} is the dipole-dipole interaction Hamiltonian between the two spins.

We will consider the spin operators \mathbf{S}_1 and \mathbf{S}_2 in the NV basis : the z orientation of the S_z operator is chosen along the NV axis of each spin. In the presence of a longitudinal magnetic field, the sense of \hat{z} will be chosen such that $\mathbf{B} \cdot \hat{z} > 0$, and the \hat{x} direction is chosen along the transverse magnetic or electric field, both of which dominate the single spin Hamiltonian. These choices mean that there will be two distinct Cartesian basis $\{\hat{x}_1, \hat{y}_1, \hat{z}_1\}$ and $\{\hat{x}_2, \hat{y}_2, \hat{z}_2\}$ describing the two spins.

The dipole-dipole Hamiltonian can then be decomposed as :

$$-\frac{\mathcal{H}_{\text{dd}}}{J_0/r^3} = 3(\mathbf{S}_1 \cdot \hat{u})(\mathbf{S}_2 \cdot \hat{u}) - \mathbf{S}_1 \cdot \mathbf{S}_2 \quad (9)$$

$$= [3(\hat{u} \cdot \hat{x}_1)(\hat{u} \cdot \hat{x}_2) - \hat{x}_1 \cdot \hat{x}_2] S_x^1 S_x^2 \quad (10)$$

$$+ [3(\hat{u} \cdot \hat{y}_1)(\hat{u} \cdot \hat{y}_2) - \hat{y}_1 \cdot \hat{y}_2] S_y^1 S_y^2 \quad (11)$$

$$+ [3(\hat{u} \cdot \hat{x}_1)(\hat{u} \cdot \hat{y}_2) - \hat{x}_1 \cdot \hat{y}_2] S_x^1 S_y^2 \quad (12)$$

$$+ [3(\hat{u} \cdot \hat{y}_1)(\hat{u} \cdot \hat{x}_2) - \hat{y}_1 \cdot \hat{x}_2] S_y^1 S_x^2 \quad (13)$$

$$+ [3(\hat{u} \cdot \hat{z}_1)(\hat{u} \cdot \hat{z}_2) - \hat{z}_1 \cdot \hat{z}_2] S_z^1 S_z^2, \quad (14)$$

$$+ \mathcal{H}_{\text{other}}, \quad (15)$$

where $J_0 = \frac{\mu_0 \gamma_e^2 \hbar^2}{4\pi} = (2\pi)52 \text{ MHz} \cdot \text{nm}^3$, $\mathbf{r} = r\hat{\mathbf{u}}$ is the distance between the two spins and $\mathcal{H}_{\text{other}}$ contains terms of the form $S_x^i S_z^j$ and $S_y^i S_z^j$ which couple states far from resonance and will be neglected here.

D. Flip-flops in the magnetic basis $\{|0\rangle, | +1\rangle, | -1\rangle\}$

We will first consider the case, treated in [2], where the longitudinal magnetic field is strong enough that the single spin Hamiltonian eigenstates of both spins are close to $\{|0\rangle, | +1\rangle, | -1\rangle\}$ (since we are interested in near-resonant spins, this means that both spins see roughly the same longitudinal and transverse magnetic field).

In this scenario, only flip-flop terms (that is $\langle \pm 1, 0 | \mathcal{H}_{\text{dd}} | 0, \pm 1 \rangle$) can couple two resonant two-spins states. Following the notation in [2], we introduce the dimensionless factor η defined as :

$$\eta^2 = \frac{1}{3} \left| \langle \pm 1, 0 | \frac{\mathcal{H}_{\text{dd}}}{J_0/r^3} | 0, \pm 1 \rangle \right|^2 \frac{4\gamma_f^2}{(\omega_f - \omega_{NV})^2 + 4\gamma_f^2}, \quad (16)$$

where γ_f is the fluctuator lifetime, $\omega_f/2\pi$ is the transition frequency of the fluctuator and $\omega_{NV}/2\pi$ the NV transition frequency.

This numerical factor is linked to the previously mentioned $\bar{\eta}$ by averaging over every possible orientation and angular position of the fluctuators. Assuming that the fluctuators are evenly distributed among all four classes of NVs and decomposing the \mathbf{r} vector in the spherical basis (r, θ, ϕ) , we can write $\bar{\eta}$ as :

$$\bar{\eta} = \frac{1}{4} \sum_{i=1}^4 \int_{\theta} \int_{\phi} \int_{\omega_f} \int_{\omega_{NV}} |\eta(\mathbf{u}, i, \omega_{NV}, \omega_f)| d\Omega \rho(\omega_{NV}) d\omega_{NV} \rho^i(\omega_f) d\omega_f, \quad (17)$$

where $d\Omega = \sin\theta d\theta d\phi$, $\rho(\omega_{NV})$ is the distribution of angular frequencies of the probed NV centers, and $\rho^i(\omega_f)$ is the distribution of angular frequencies for the fluctuators of class i .

While the distributions $\rho(\omega_{NV})$ and $\rho^i(\omega_f)$ can be approximated using the data presented in Fig. 6, we will assume that these distributions are independent of the external magnetic field, and since we are interested in comparing the same sample for various values of the magnetic field, we will simplify eq.(17) by setting $\frac{4\gamma_f^2}{(\omega_f - \omega_{NV})^2 + 4\gamma_f^2}$ to 1 if the classes of NV centers and fluctuators are resonant (or if they are from the same class), and to 0 otherwise.

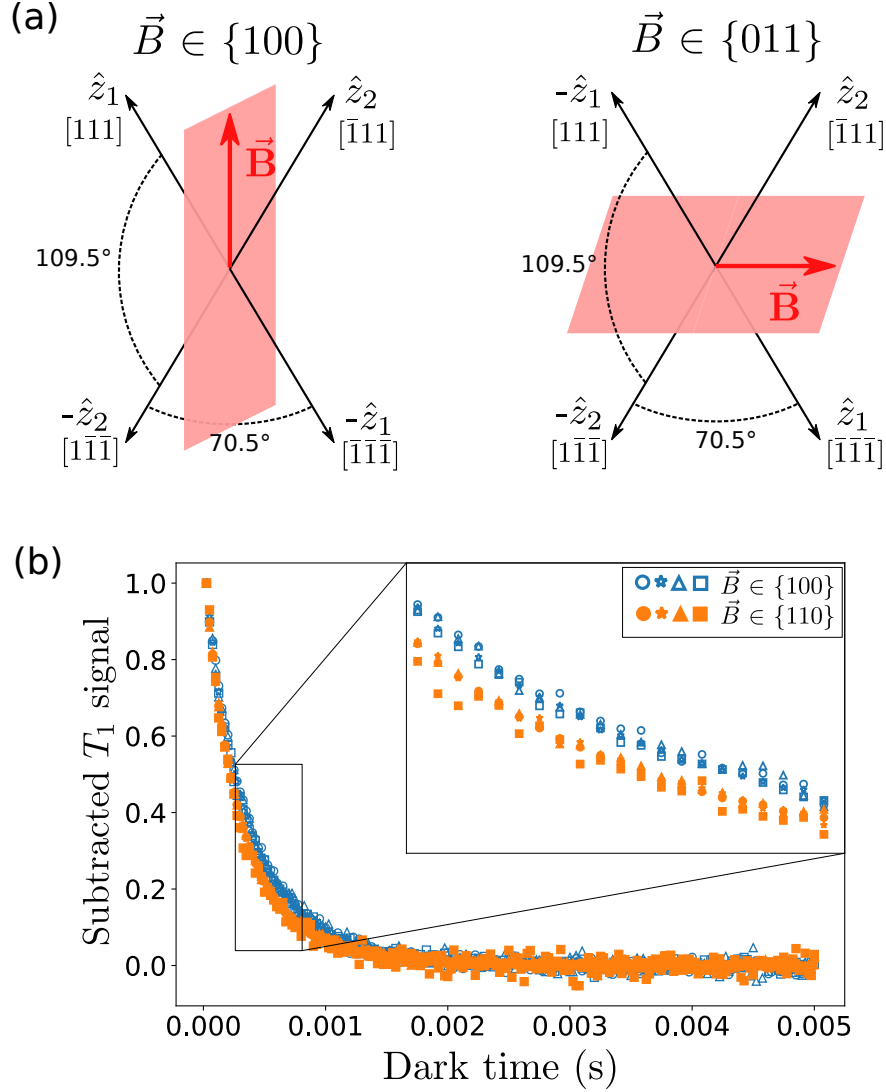


FIG. 12. (a) Geometrical representation of two NV classes and the two possible planes where the magnetic field has the same projection on both classes. The positive \hat{z}_i direction is chosen so that $\vec{B} \cdot \hat{z}_i > 0$. (b) T_1 measurement on a two-class resonance following the protocol described in main text. 8 different magnetic field were employed on the same sample : 4 times in a $\{100\}$ plane (blue unfilled symbols) and 4 times in a $\{110\}$ plane (orange filled symbols). Inset : zoom-in on the 0.3-0.8 ms region.

With these hypotheses, we find that there are 3 possible scenarios where the NV and fluctuator are resonant :

- The NV and fluctuator are from the same class, so that the angle $\widehat{z_1 z_2} = 0$: this is always true, regardless of the magnetic field angle and amplitude. We can analytically

compute $\bar{\eta}$ in this case and find :

$$\bar{\eta}_{same} = \frac{1}{4} \int_{\theta} \int_{\phi} |\eta(\mathbf{u}, \widehat{z_1 z_2} = 0, \omega_{NV} = \omega_f)| d\Omega = \frac{1}{4} \cdot \sqrt{\frac{1}{3}} \cdot \frac{2}{3\sqrt{3}} \approx 5.55 \cdot 10^{-2}.$$

- The angle $\widehat{z_1 z_2} = \arccos(\frac{1}{3}) \approx 70.5^\circ$. This is the case when the magnetic field lies in the $\{100\}$ crystalline planes family. In this case, numerical simulations yield $\bar{\eta}_{close} \approx \frac{1}{4} \cdot \sqrt{\frac{1}{3}} \cdot 0.6507 \approx 9.39 \cdot 10^{-2}$
- The angle $\widehat{z_1 z_2} = \arccos(-\frac{1}{3}) \approx 109.5^\circ$. This happens when the magnetic field lies in the $\{110\}$ or $\{1\bar{1}0\}$ crystalline planes family. In this case, numerical simulations give $\bar{\eta}_{far} \approx \frac{1}{4} \cdot \sqrt{\frac{1}{3}} \cdot 0.8328 \approx 1.20 \cdot 10^{-1}$. This last case was not taken into account in [2].

Fig. 12(a) shows a graphical representation of the difference between the two last scenarios : due to the $\mathbf{B} \cdot \hat{\mathbf{z}} > 0$ condition, the same two classes of NV centers can have a $\widehat{z_1 z_2}$ angle equal to 70.5° or 109.5° depending on the external magnetic field.

These computed values can be tested experimentally. Fig. 12(b) shows the subtracted T_1 signal measured on the same sample HPHT-15-2 for 8 different values of the magnetic field, each time on a two-class resonance. In 4 cases, the magnetic field was in a $\{100\}$ plane, and in the 4 other cases it was in a $\{110\}$ plane. We can see that the measured lifetime is always smaller on the $\{110\}$ case, which corresponds to the greater $\bar{\eta}$ factor computed previously.

Here is a list of the computed values of $\bar{\eta}^2$, which according to eq.(6) is proportional to the dipole induced decay rate, for various magnetic field orientation. An ODMR spectrum for each of these situations is present in Fig. 13.

- Random orientation / no class degeneracy : for all four classes, $\bar{\eta} = \bar{\eta}_{same}$ and $\bar{\eta}^2 = 3.08 \cdot 10^{-3} \equiv \bar{\eta}_0^2$.
- $\mathbf{B} \in \{110\}$: For the two non-resonant classes : $\bar{\eta} = \bar{\eta}_{same}$ and $\bar{\eta}^2 = \bar{\eta}_0^2$. For the two resonant classes : $\bar{\eta} = \bar{\eta}_{same} + \bar{\eta}_{far}$ and $\bar{\eta}^2 \approx 10.0 \bar{\eta}_0^2$.
- $\mathbf{B} \in \{100\}$: For each pair of two-classes resonance: $\bar{\eta} = \bar{\eta}_{same} + \bar{\eta}_{close}$ and $\bar{\eta}^2 \approx 7.24 \bar{\eta}_0^2$.
- $\mathbf{B} \parallel \langle 111 \rangle$: For the non-resonant class (parallel to \mathbf{B}) : $\bar{\eta} = \bar{\eta}_{same}$ and $\bar{\eta}^2 = \bar{\eta}_0^2$. For the triply resonant classes: $\bar{\eta} = \bar{\eta}_{same} + 2\bar{\eta}_{far}$ and $\bar{\eta}^2 \approx 28.4 \bar{\eta}_0^2$.
- $\mathbf{B} \parallel \langle 100 \rangle$: For the quadruple resonance : $\bar{\eta} = \bar{\eta}_{same} + 2\bar{\eta}_{close} + \bar{\eta}_{far}$ and $\bar{\eta}^2 \approx 42.8 \bar{\eta}_0^2$.

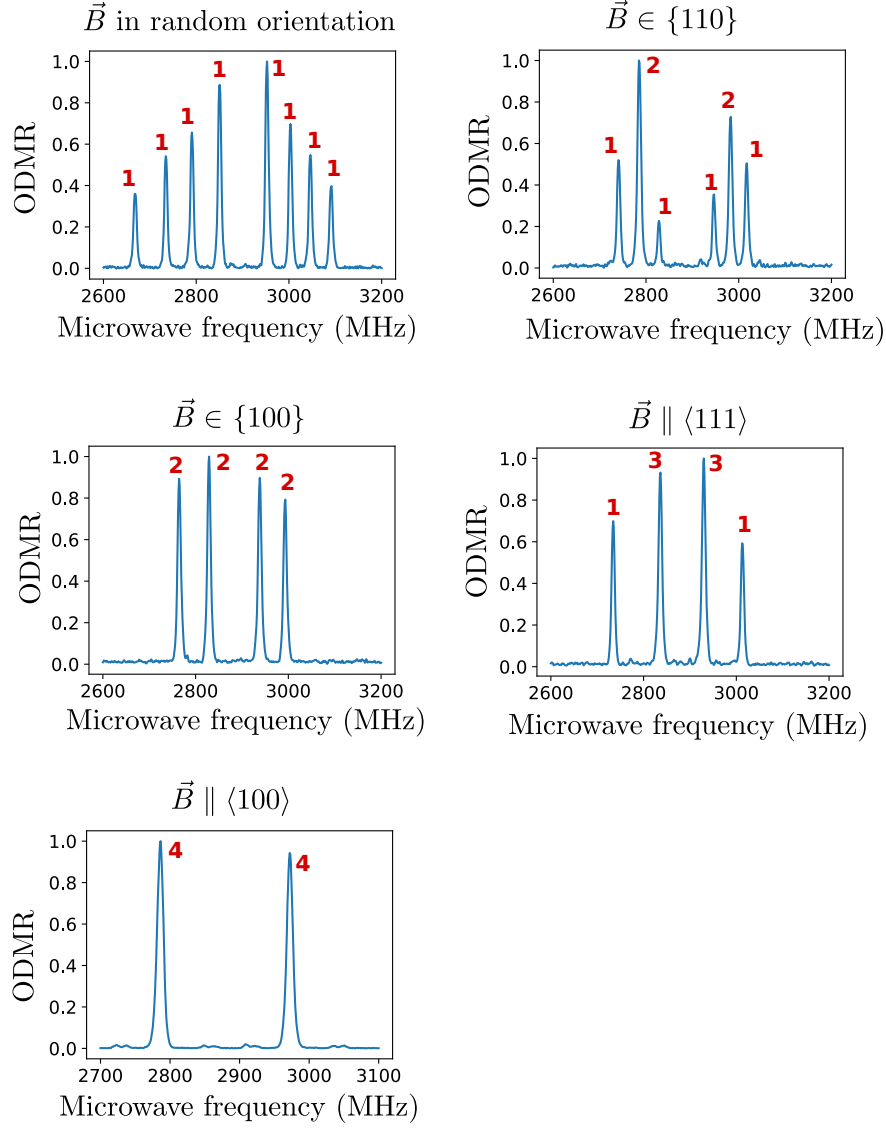


FIG. 13. ODMR spectra for different orientations of the magnetic field. Red numbers represent the number of degenerate classes for each line of the spectrum.

The measured increase in the decay rate is generally smaller than the predicted one : we measured an increase by a factor ~ 4 for a two-class degeneracy instead of the predicted 10.0, and an increase by a factor ~ 16 for a four-class degeneracy instead of the predicted 42.8. [2] measured an increase by a factor ~ 4 on a two-class degeneracy instead of the predicted 7.24 or 10.0.

E. Flip-flops in the non-magnetic basis $\{|0\rangle, |+\rangle, |-\rangle\}$

As described in sec. I, when the single spin Hamiltonian of the NV center is dominated by the electric field or the transverse magnetic field (as long as $\gamma_e B_\perp \ll D$), the eigenbasis of the NV Hamiltonian is close to $\{|0\rangle, |+\rangle = \frac{|+1\rangle + |-1\rangle}{\sqrt{2}}, |-\rangle = \frac{|+1\rangle - |-1\rangle}{\sqrt{2}}\}$. We therefore have to rewrite the dipole-dipole Hamiltonian using this new basis for both spins in order to perform similar calculations as the ones done in the previous section.

Since the flip-flop processes are now of the form $|0, \pm\rangle \langle \pm, 0|$, we redefine the η factor as :

$$\eta^2 = \frac{1}{3} \left| \langle \pm, 0 | \frac{\mathcal{H}_{\text{dd}}}{J_0/r^3} | 0, \pm \rangle \right|^2 \frac{4\gamma_f^2}{(\omega_f - \omega_{NV})^2 + 4\gamma_f^2}. \quad (18)$$

In this new basis, the spin operators are written :

$$S_x = \begin{pmatrix} 0 & 1 & 0 \\ 1 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix} \begin{matrix} \langle -| \\ \langle 0| \\ \langle +| \end{matrix}, \quad S_y = \begin{pmatrix} 0 & 0 & 0 \\ 0 & 0 & 1 \\ 0 & 1 & 0 \end{pmatrix} \begin{matrix} \langle -| \\ \langle 0| \\ \langle +| \end{matrix}, \quad S_z = \begin{pmatrix} 0 & 0 & 1 \\ 0 & 0 & 0 \\ 1 & 0 & 0 \end{pmatrix} \begin{matrix} \langle -| \\ \langle 0| \\ \langle +| \end{matrix}. \quad (19)$$

The symmetry in the (xy) plane is broken by the presence of the transverse electric or magnetic field. The x_i direction is defined as the direction of the transverse electric or magnetic field which dominates the single spin Hamiltonian of the particle i .

We will consider two cases :

- The x direction is defined by an external field (transverse magnetic field or strong external electric field) projected on the (xy) plane of each classes. In particular this means that two spins from the same class have the same $(\hat{x}, \hat{y}, \hat{z})$ basis.
- The x direction is defined by the local electric field generated by the charges in the crystal. We will assume that the NV and fluctuator do not see the same electric field, and will sample a random angle $\psi \in [0, 2\pi]$ between the axes \hat{x}_1 and \hat{x}_2 .

For this two cases, which we will call $\hat{x}_1 = \hat{x}_2$ and $\hat{x}_1 \neq \hat{x}_2$, we can compute the $\bar{\eta}$ factor for the three scenarios discussed previously : $\widehat{z_1 z_2} = 0$ (same class), $\widehat{z_1 z_2} = 70.5^\circ$ and $\widehat{z_1 z_2} = 109.5^\circ$. The results, as well as those in the $|\pm 1\rangle$ basis are presented in Table I.

There are two particular situations where these values can be experimentally tested :

The first one is described in Fig. 4 of the main text : in the presence of pure transverse magnetic field (for a single class, non-resonant with the three other classes), the decay rate

$\bar{\eta}$ table	$\widehat{z_1 z_2} = 0$	$\widehat{z_1 z_2} = 70.5^\circ$	$\widehat{z_1 z_2} = 109.5^\circ$
$ \pm 1\rangle$ basis	$\frac{2}{3\sqrt{3}} = 0.3849$	0.6507	0.8328
$ +/-\rangle$ basis $\hat{x}_1 \neq \hat{x}_2$	0.7110	0.6828	0.6828
$ +/-\rangle$ basis $\hat{x}_1 = \hat{x}_2$	$\frac{4}{3\sqrt{3}} = 0.7698$	0.6951	0.6951

TABLE I. Computation of $\frac{\bar{\eta}}{\frac{1}{4}\sqrt{\frac{1}{3}}}$ for the different eigenbasis of the single spin Hamiltonian, and for the different angles between the z axis of the two spins.

of the class decreases with the magnetic field amplitude, due to the double-flip processes (described below), and then reaches a plateau with a decay rate value ~ 2 times larger than in the longitudinal magnetic field case. This increase is in agreement with the fact that $\bar{\eta}_{\text{same}}$ is 2 times bigger in the non-magnetic $|+/-\rangle$ basis than in the magnetic $|\pm 1\rangle$ basis. The measured increase is again smaller than the predicted factor of 4 given by eq. (6).

The second one is the difference between the $\mathbf{B} \parallel \langle 100 \rangle$ case and the $\mathbf{B} = 0$ case. When a magnetic field is applied in the $[100]$ direction, all four classes are resonant and the spin Hamiltonian basis is $\{|0\rangle, | +1 \rangle, | -1 \rangle\}$. The $\bar{\eta}$ factor in this case has been calculated in the previous section : $\bar{\eta}^2 \approx 42.8 \bar{\eta}_0^2$. When no external magnetic field is applied, all four classes are also resonant but the spin Hamiltonian basis is $\{|0\rangle, | + \rangle, | - \rangle\}$ (see Sec. 1) and double-flip processes are near-resonant. The decay rate due purely to flip-flop in the $|+/-\rangle$ basis is proportional to $\bar{\eta}^2 = (\bar{\eta}_{\text{same}}^{+/-} + 3\bar{\eta}_{\text{diff}}^{+/-})^2 \approx 51.4 \bar{\eta}_0^2$. Fig. 3 of the main text shows that we indeed observe a shorter lifetime when $\mathbf{B} = 0$, in agreement with the higher value of $\bar{\eta}^2$, but this decrease could also come from the double-flip processes which are absent when $\mathbf{B} \neq 0$.

F. Double-flip processes

Double-flip processes are processes where both spins lose or gain one unit of spin angular momentum, as opposed to flip-flop processes where one spin loses one quantum and the other gains one. These are related to matrix elements such as $\langle +1, 0 | \frac{\mathcal{H}_{\text{dd}}}{J_0/r^3} | 0, -1 \rangle$ in the $|\pm 1\rangle$ basis or $\langle +, 0 | \frac{\mathcal{H}_{\text{dd}}}{J_0/r^3} | 0, - \rangle$ in the $|\pm\rangle$ basis. These matrix elements couple two-spin states that are never fully resonant, however with low to zero magnetic, the residual splitting due to local electric and magnetic field (~ 6 MHz with our samples) is small enough compared to the fluctuators linewidth measured in sec. IV B to be ~ 8 MHz, so that double-flip processes may still occur.

Fig. 4 of the main text shows that for lower transverse field values, the spin decay rate is increased significantly. We attribute this increase to double-flip processes. We can see that the decay rate increase (~ 5 times the baseline value) is significantly higher than the increase due solely to the $|+/-\rangle$ basis (~ 2 times the baseline value). This lead us to believe that the double-flip process are the main reason behind the decrease in spin lifetime for $\mathbf{B} = 0$ observed in Fig. 4 of the main text. This would probably mean that double-flip processes are the dominant factor in the sensitivity of the magnetometry protocol presented in the main text.

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