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Laser threshold magnetometry using green-light absorption by diamond nitrogen vacancies in an external cavity laser

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Nitrogen vacancy (NV) centers in diamond have attracted considerable recent interest for use in quantum sensing, promising increased sensitivity for applications ranging from geophysics to biomedicine. Conventional sensing schemes involve monitoring the change in red fluorescence from the NV center under green laser and microwave illumination. Due to the strong fluorescence background from emission in the NV triplet state and low relative contrast of any change in output, sensitivity is severely restricted by a high optical shot noise level. Here, we propose a means to avoid this issue, by using the change in green pump absorption through the diamond as part of a semiconductor external cavity laser run close to the lasing threshold. We show that theoretical sensitivity to the magnetic field on the $\text{pT}/\sqrt{\text{Hz}}$ level is possible using a diamond with an optimal density of NV centers. We discuss the physical requirements and limitations of the method, particularly the role of amplified spontaneous emission near threshold and explore realistic implementations using current technology.

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I. INTRODUCTION

Optical manipulation of material defects represents an ideal method for quantum sensing, exploiting properties such as entanglement and superposition [1]. The nitrogen-vacancy (NV) center in diamond, possessing long quantum coherence times at room temperature, has in particular drawn considerable interest [2–4]. Diamond is an ideal material for sensing, being mechanically hard, chemically stable, isotopically pure, as well as biocompatible [5,6]. The negatively charged nitrogen-vacancy center (NV^-) has an energy level structure that results in optical properties that are highly sensitive to temperature [7], strain (pressure) [8], electric field [9], and, particularly, magnetic field. Sensing is conventionally performed by detecting changes in the intensity of red fluorescence ($\approx 637\text{--}750\text{ nm}$) under irradiation with green light and resonant microwaves via a process termed optically detected magnetic resonance (ODMR) spectroscopy [4,10–12]. It can be done using a continuous wave (cw) method [13] or by using short laser and microwave pulses [14,15].

However, measuring via red fluorescence suffers from two considerable physical limitations. First, the signal to be measured has a very low contrast on bright emission from decay in the NV^- triplet state. For a single NV^- or for a smaller ensemble where the NV centers are preferentially aligned through growth engineering [16], spin-dependent contrast can be up to 30% [17]. However, for bulk sensing using an ensemble of many billions of NV centers in a region encompassing a large volume of the diamond sensor (up to the whole diamond), the

contrast that is realized is often, at most, a few percent [18,19]. The sensitivity is therefore limited by this low contrast and the high level of shot noise from the bright background rising from triplet state fluorescence emission. The second physical limitation is the high refractive index of diamond, which traps the majority of the fluorescence inside the diamond. Micro-fabrication schemes have been proposed to mitigate this issue, but have yet to deliver significant improvements [20,21].

An alternative method is to use optical absorption of the pump light by the NVs. Previous work has used the change in green absorption in an optical cavity [22,23] or by using changes in infrared (IR) absorption by the singlet state [24]. These schemes are technically demanding, requiring an optical cavity or unusual wavelength (1042 nm) laser. A promising alternative is laser threshold sensing [25], using changes in optical absorption resulting from the parameter to be sensed (e.g., magnetic field or temperature) to push a medium across the lasing threshold. This method eliminates the bright background that limits sensitivity using conventional fluorescence detection. A further attraction is the wide applicability to any material with variable optical absorption, including a wider range of defects in diamond, SiC, and two-dimensional (2D) materials [26,27].

Building on the work by Dumeige *et al.* [28] and our own previous work on diamond absorption magnetometry [22,23], here we outline a scheme to use laser threshold sensing of the magnetic field with green light in a standard external cavity laser. We show it is possible to achieve high sensitivity in the $\text{pT}/\sqrt{\text{Hz}}$ range with realistic assumptions for the key physical parameters. Our proposal differs from that of previous work by using simpler green pump absorption rather than IR absorption and by using an ordinary current driven laser

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FIG. 1. (a) Schematic of the external cavity setup with a Fabry-Perot semiconductor laser diode of cavity length L_m and end facet power reflectivities R_1 and R_2 coupled to an external cavity of length L_r via mirror R_3 and diamond of thickness d . Laser emission (dashed line) is through R_1 . We model our external cavity laser with the diamond as a single cavity with equivalent end reflectance R_e and the diamond absorption loss α_d included in the total cavity loss α_t . (b) Simplified schematic of the laser threshold process, where a reduction in diamond absorption through application of resonant microwaves reduces the threshold current to I_{th}^{on} , producing lasing output P_{out} when driven at I_{th}^{off} .

diode or gain chip medium without the need for an additional pump laser. We show that this configuration, which is highly suitable for miniaturization, can deliver high sensitivity, and we discuss the key physics required to reach such sensitivity levels. Finally, we discuss and calculate limiting factors that may prevent these levels from being reached in practice. This includes factors that may not have been previously considered, such as amplified spontaneous emission near to the lasing threshold.

II. METHODS

A. External cavity laser model

We place the diamond into a standard external cavity laser setup as described schematically in Fig. 1(a). This consists of a Fabry-Perot semiconductor laser diode or gain chip of length L with end facet reflectivities R_1 and R_2 coupled to an external cavity formed by mirror R_3 via an external cavity of length L_r containing the diamond of thickness d . We assume normal incidence and that transmission through the diamond

is high, with minimal reflection from the diamond facets. For simplicity, we assume a single optical mode at a single wavelength. We consider the optical loss due to absorption in a diamond of thickness d . The change in laser intensity $\hat{I}$ on a pass through the diamond is given by

$$\Delta \hat{I} = \hat{I}_0 - \hat{I}_0 e^{-\alpha_d z}, \quad (1)$$

where $\hat{I}_0$ is the intensity of the laser emission with no diamond present in the external cavity, α_d is the absorption coefficient in the diamond, and z is the path length taken within the diamond. For normal incidence, $z = d$ and the absorption coefficient can be derived from the rate equation model given in the following section or can be measured experimentally. For the semiconductor lasing medium between mirrors 1 and 2, we assume a total cavity loss α_t , given by the sum of intrinsic cavity loss due to the gain medium α_c and losses from the mirrors and end facets α_m , giving a total loss α_t ,

$$\alpha_t = \alpha_c + \alpha_m = \alpha_c + \frac{1}{L_m} \ln \left(\frac{1}{\sqrt{R_1 R_2}} \right). \quad (2)$$

In order to simplify the analysis of the external cavity structure, we use the three mirror model [29–32] to treat the complete diode and external cavity structure as a single cavity of length $L = L_m + L_r$, with mirror R_2 replaced by an effective reflectivity R_e , with the single cavity containing the optical losses of the external cavity and diamond, the internal losses of the gain medium in the laser diode, and the loss from the cavity through the mirrors. By assuming that the losses due to the diamond are spread evenly throughout, we redefine the loss coefficient due to the diamond as $\alpha_e = (\alpha_d/L)d$, and our total cavity loss as

$$\alpha_t = \alpha_c + \alpha_e + \frac{1}{L} \ln \left(\frac{1}{\sqrt{R_1 R_e}} \right), \quad (3)$$

where $R_e = |r_e|^2$ relates the power reflectivity to the complex field reflectivity r_e . We use the model for the effective reflectivity by Voumard *et al.* [33], detailed further in the Supplemental Material [34]. Neglecting phase components, at threshold, $R_1 R_e e^{(\Gamma g - \alpha_t)2L} = 1$, where $g = g_{th}$ is the (threshold) gain coefficient.

For the full structure, the rate equations for photon (S) and carrier (N) density are given by the standard equations for a laser diode as

$$\frac{dN}{dt} = \frac{I}{qV} - \frac{N}{\tau_N} - GS \quad (4)$$

and

$$\frac{dS}{dt} = GS - \frac{S}{\tau_p} + \frac{\beta N}{\tau_N}. \quad (5)$$

Here, I is the drive current, V the volume of the gain region, G the gain of the lasing medium, and q the electronic charge. The term GS arises from stimulated emission in the laser diode gain medium and S/τ_p includes the cavity loss from the mirrors, gain medium, and diamond. Further, τ_p is the photon lifetime in the cavity and τ_N the carrier lifetime in the laser diode. Carriers are generated by a current I in a volume V , where $V = L \times w \times t_h$, where t_h is the thickness and w the width of the laser diode active region. The term $\beta N/\tau_N$

relates to spontaneous emission, governed by the spontaneous emission factor β .

We can define gain G phenomenologically, in the form [35]

$$G = \Gamma g = \Gamma a(N_{th} - N_{tr})(1 - \epsilon S), \quad (6)$$

where Γ is the confinement factor and ϵ is the gain compression factor that phenomenologically accounts for effects such as spectral hole burning at higher optical power. The carrier density at transparency is given by N_{tr} . The rate equations for photon and carrier density can be solved for a steady state condition ($dS/dt = 0$, $dN/dt = 0$). For carrier density $N = N_{th}$ close to N_{tr} and neglecting spontaneous emission ($\beta = 0$), the gain balances the cavity loss. The factor a is the differential gain coefficient, a material specific property defining how well the semiconductor can generate carriers for population inversion. Equation (6) is valid for heterostructure laser diodes and certain quantum well structures where the threshold is close to the transparency carrier density. Unless otherwise stated, we use the model of Eq. (6) in this work.

Using Eqs. (4)–(6) at lasing threshold, where $S = 0$, $G = 1/\tau_p$, and $\Gamma g_{th} = \alpha_t$, we can derive an equation for carrier density at threshold N_{th} ,

$$N_{th} = N_{tr} + \frac{\alpha_t}{\Gamma a}, \quad (7)$$

and inserting this result into the rate equation for carrier density [Eq. (4)] allows us to calculate the threshold current,

$$I_{th} = \frac{qV}{\eta_i \tau_N} N_{th} = \frac{qV}{\eta_i \tau_N} \left(N_{tr} + \frac{\alpha_t}{\Gamma a} \right). \quad (8)$$

Here we introduce the quantum efficiency of the carrier to photon conversion η_i . By using Eq. (8) in the rate equations at $I > I_{th}$, we can calculate the photon density at any current above the lasing threshold. We can then calculate the laser light power that can be emitted from the left-hand side mirror R_1 using the factor η_o , the output coupling efficiency, which is defined as the ratio of photons lost through the mirror R_1 to the total cavity loss $\alpha_t = \alpha_m + \alpha_c + \alpha_e$,

$$P_{out} = \eta_o \frac{hc}{\lambda \tau_p} \frac{V}{\Gamma} S, \quad (9)$$

where V/Γ is the effective mode volume of the cavity, λ the wavelength, and h and c Planck's constant and the speed of light, respectively. In the limit of $\epsilon S \rightarrow 0$ where there is no limiting effect on the gain, the power output can be rewritten directly in terms of the threshold current,

$$P_{out} = \eta_o \eta_i \frac{hc}{q\lambda} (I - I_{th}). \quad (10)$$

In both of these expressions,

$$\eta_o = \frac{\alpha_{m1}}{\alpha_m + \alpha_c + \alpha_e} = \frac{\ln \frac{1}{\sqrt{R_1}}}{\ln \frac{1}{\sqrt{R_1 R_e}} + (\alpha_e + \alpha_c)L}. \quad (11)$$

For larger finite values of ϵ well above threshold or including finite spontaneous emission through nonzero β , we can numerically solve the steady state rate equations [Eqs. (4) and (5)] to calculate N , S , and the laser power output.

The total cavity absorption α_t will change when microwaves are applied to the diamond at a frequency equal to

the splitting of the NV triplet ground state levels, reducing the lasing threshold current, $\Delta I_{th} = I_{th}^{off} - I_{th}^{on}$, where I_{th}^{on} is the threshold current *on* microwave resonance and I_{th}^{off} the threshold current *off* resonance. By running at drive current equal to I_{th}^{off} , the laser output is generated only while on microwave resonance. This is shown schematically in Fig. 1(b).

B. Absorption model

We use the rate equation model from [36] in order to calculate the optical absorption of green pump light by the diamond and the maximum change in absorption when on microwave resonance. The parameters we use for the transition rates are the same as those in [28], derived from [37–40]. We calculate the normalized occupancies of each energy level with microwaves supplied n_i^{on} and without microwaves n_i^{off} , where $\sum_i n_i = 1$ and index $i = 1-8$, where $i = 1$ refers to the $m_s = 0$ ground state level, $i = 2$ the $m_s = \pm 1$ ground state levels, $i = 3, 4$ the spin triplet excited states, $i = 5, 6$ the spin singlet shelving states, and $i = 7, 8$ the ground and excited state of the NV^0 . We define a total NV^- density N_{NV} in ppm. Off resonance, the total number density of NV^- in each state N_i^{off} is given by

$$N_i^{off} = N_{NV} \frac{n_i^{off}}{\sum_i n_i^{off}}. \quad (12)$$

We define a measurement axis along one of the four possible crystallographic axes for the NV. We calculate that when microwaves are applied, we drive only the NVs aligned along one axis such that the total number density on resonance N_i^{on} is given by

$$N_i^{on} = \frac{1}{4} N_{NV} \frac{n_i^{on}}{\sum_i n_i^{on}} + \frac{3}{4} N_{NV} \frac{n_i^{off}}{\sum_i n_i^{off}}. \quad (13)$$

We calculate the change in intensity on a single pass when on and off microwave resonance as

$$\hat{I}_{on} = \hat{I}_0 e^{-\alpha_{on} d}, \quad \hat{I}_{off} = \hat{I}_0 e^{-\alpha_{off} d}, \quad (14)$$

where d is the thickness of the diamond, and the absorption coefficient α on and off resonance is given by

$$\alpha^{on} = \sigma_g(N_1^{on} + N_2^{on}) + \sigma_{g0}N_7^{on} + \sigma_e(N_3^{on} + N_4^{on}) + \sigma_rN_8^{on}, \quad (15)$$

$$\alpha^{off} = \sigma_g(N_1^{off} + N_2^{off}) + \sigma_{g0}N_7^{off} + \sigma_e(N_3^{off} + N_4^{off}) + \sigma_rN_8^{off}. \quad (16)$$

Here, σ_g and σ_{g0} are, respectively, the absorption cross sections of green light for NV^- and NV^0 , and σ_e and σ_r are the ionization cross sections for transfer between the charged and uncharged defect states. This allows us to calculate the change in absorption when the diamond is present without microwaves, $\hat{I}_{off}/\hat{I}_0$, the change when driven on microwave resonance, $\hat{I}_{on}/\hat{I}_0$, and the change between these, which we term the absorption contrast,

$$C = (\hat{I}_{off}/\hat{I}_0) - (\hat{I}_{on}/\hat{I}_0). \quad (17)$$

C. Key physical parameters

The key physical parameters of the model can be divided into those that are intrinsic to the semiconductor gain medium, those intrinsic to the diamond, and those defined by the setup. Examples of the latter include the mirror reflectivities R_1 , R_2 , R_3 , the cavity length L , and any other losses, such as reflection out of the cavity or from absorption by other optical components such as lenses, included in the cavity loss factor α_c . These factors will also influence the photon lifetime in the cavity τ_p . The maximum Rabi frequency Ω_R that can be reached also depends on the microwave power and how well the microwaves can be coupled into the diamond.

The parameters which are intrinsic to the diamond are the diamond thickness d , NV^- density N_{NV} , ensemble dephasing time T_2^* defining the ODMR linewidth, and absorption contrast C arising from changes in pump absorption on or off microwave resonance. These factors define the diamond absorption factor α_d .

A number of these parameters are interrelated. The ODMR linewidth is proportional to the inverse of T_2^* , which in turn is dependent on N_{NV} concentration in the limit of high nitrogen content and the abundance of ^{13}C for low nitrogen content [41]. There is also a dependence on other material properties such as strain [42], which makes the relationship between the parameters difficult to determine. We therefore consider values in the experimental literature as a guide. Figure 2 shows a plot of T_2^* versus NV^- density N_{NV} for a range of diamonds from the literature [16,38,43–47]. Typical NV^- densities range from 10^{-4} ppm up to tens of ppm [48]. In general, $T_2^* < 1 \mu\text{s}$ for samples with natural (1.1%) ^{13}C content [49,50]. Experiments typically realize Rabi frequencies Ω_R of 1–5 MHz, with up to 10 MHz using optimal antenna geometries [51].

We include only the uncharged NV^0 and negatively charged NV^- nitrogen vacancy centers in our model and we do not consider the role of other types of defects in or on the diamond, including substitutional nitrogen or other materials that may be present (e.g., Si, boron). Furthermore, we assume a uniform distribution of nitrogen vacancies throughout the diamond, such that we can consider a single, fixed absorption cross section throughout. These are simplifications introduced due to the poorly known absorption cross sections of certain types of defects or dopants and the experimental difficulty in quantifying density within (or on) diamonds. We highlight that this implies that the sensitivities estimated by our model represent a best-case scenario, with no optical loss from the NV^- system used for sensing due to the presence of other defects, dopants, or adsorbents in or on the diamond.

Those parameters, intrinsic to the laser diode or gain chip that is used, are the carrier density at transparency N_{tr} , the gain compression factor ϵ that arises from effects that limit the gain well above threshold, the differential gain coefficient a that relates the gain and carrier density, the threshold carrier lifetime τ_N , the confinement factor Γ , the volume of the gain medium V , and the spontaneous emission factor β . For our gain medium, we take a III-V semiconductor heterostructure device, such as the nitride compounds capable of emission at green wavelengths (e.g., InGaN) [66]. Table I shows a typical range of values for each of these parameters. We

FIG. 2. Dephasing time T_2^* vs NV^- density N_{NV} , where both values are given in other works (citations given in the main text). T_2^* in those with low NV^- concentration are limited by interaction with ^{13}C spin, with the highest values given by diamonds isotopically purified with ^{12}C spin during growth. T_2^* in those with high NV^- concentration are limited by dipolar interaction between defects, including other substitutional nitrogen defects such as P1 centers. Note that the NV^- density for the work by Childress *et al.* is an upper estimate made here assuming a 10% NV^- fraction; total substitutional nitrogen content for this diamond was given as ≤ 0.1 ppm. (Both axes use a $\log_{10}$ scale.)

take the typical ranges shown based on experimental results from different structures (quantum well, vertical cavity) and from calculations based on bulk material properties such as effective mass. N_{tr} effectively defines the size of the lasing threshold current I_{th} . The desired change in threshold current on change in absorption factor α_t is defined, in particular, by Γ and the gain coefficient a in Eqs. (3) and (7).

TABLE I. Typical ranges for the key semiconductor gain medium parameters. Here, N_{tr} , τ_N , and a are taken for typical III-nitride semiconductors. The range for Γ is given for laser diodes with a thin (sub- μm) active layer and is typically no more than a few percent. The range of β is given for the literature values for a range of laser diodes where confinement is not deliberately sought, e.g., microcavities, and where values several orders of magnitude higher than the given range are possible [65].

Parameter	Range	Ref.
Transp. carrier density, N_{tr}	$3 \times 10^{18} - 2 \times 10^{19} \text{ cm}^{-3}$	[52–56]
Carrier lifetime, τ_N	1–5 ns	[57,58]
Differential gain factor, a	$10^{-17} - 10^{-22} \text{ m}^2$	[59,60]
Confinement factor, Γ	0.01–0.1	[61,62]
Spont. emission factor, β	$10^{-5} - 10^{-2}$	[63,64]

FIG. 3. (a) Absorption contrast percentage calculated from the rate model for diamond D3. This is the maximum change in absorption between on microwave resonance and off microwave resonance as a function of Rabi frequency Ω_R and laser intensity $\hat{I}$ in W/m^2 . (b)–(d) Normalized level occupancy for the NV^- triplet ground state, responsible for green absorption, the uncharged NV^0 defect state, and the NV^- singlet state. At high laser intensities, population transfer to NV^0 limits the achievable absorption contrast. For reference, 10 mW of laser power with a 1-mm-diameter circular beam on the diamond gives an intensity $\hat{I} = 10^4 \text{ W}/\text{m}^2$. The black spot indicates the Rabi frequency and power for calculations later in this work. (All x and y axes use a $\log_{10}$ scale.)

III. RESULTS

A. Absorption contrast

We first calculate, from the rate model, the fraction of incident pump light which is absorbed by the diamond and the change in this absorption (C) when on microwave resonance. We choose to model three different diamonds covering different regimes: D1, D2, and D3 with parameters (NV density and T_2^*) representative of the values seen in the literature (Fig. 2). For diamond D1, we choose a low NV^- concentration $N_{\text{NV}} = 0.001 \text{ ppm}$ and high $T_2^* = 5 \mu\text{s}$, representative of ^{12}C enriched diamonds. For diamond D2, we choose a medium NV^- concentration $N_{\text{NV}} = 0.1 \text{ ppm}$ and $T_2^* = 0.75 \mu\text{s}$, representative of chemical vapor deposition (CVD)-grown diamond with natural ^{13}C abundance. For diamond D3, we choose $N_{\text{NV}} = 10 \text{ ppm}$ and $T_2^* = 0.1 \mu\text{s}$, characteristic of high nitrogen content high-pressure high-temperature (HPHT) diamond. We use a diamond thickness $d = 500 \mu\text{m}$ for all, representative of commercially available single-crystal plates.

Using the rate model, we can calculate the absorption of light incident on the diamond D1: $\hat{I}_{\text{off}}/\hat{I}_0$, where $\hat{I}_0$ is the intensity of the incident light and $\hat{I}_{\text{off}}$ is the intensity of the light after the diamond (without supplying microwaves). The calculated absorption for diamonds D1–D3 is 0.015%, 1.498%, and 77%, respectively, as expected from increasing NV density. We can also calculate the change in absorption when on and off microwave resonance. This is shown in Fig. 3 as absorption contrast C for D3 as a function of microwave drive power (as Rabi frequency Ω_R) and laser output (as intensity). The equivalent plots for D1 and D2 are given in the Supplemental Material [34]. The maximum $C = 0.22\%$ for D3 and lowest

for D1 with the lowest NV density with $C = 10^{-4}\%$. This contrast is comparable to our previous absorption experiments using a diamond with equivalent ppb-level NV^- density [22]. We note that at Rabi frequencies above 100 kHz and laser outputs above $10^6 \text{ W}/\text{m}^2$, the absorption contrast begins to drop. This results from depopulation of the triplet ground state $^3\text{A}_2$ [normalized occupancy in Fig. 3(b)] in favor of the NV^0 [Fig. 3(c)] and the singlet shelving state [Fig. 3(d)]. However, since we aim to operate near the lasing threshold, laser intensity will be low in our scheme, avoiding this issue and ensuring that we remain in the region of highest contrast.

To further validate the absorption modeling, we have also measured diamond absorption on a high-density sample consisting of a 1-mm-thick HPHT diamond with 200 ppm nitrogen content, irradiated with 10 MeV electrons and annealed at 900°C . The estimated NV content for this sample was 10–20 ppm. The absorption contrast for this sample is shown in Fig. 4. The sample was found to be moderately polycrystalline and was therefore measured without an offset field to produce a single central dip in fluorescence, with a number of satellite features resulting from the polycrystallinity and residual magnetic field in the laboratory. Here, total off-resonance diamond absorption was 90% of incident pump light and maximum absorption contrast $C = 0.13\%$. For comparison to experiment, we model absorption contrast with our model with NV density of 15 ppm and $T_2^* = 100 \text{ ns}$, derived from an estimate of the resonance linewidth, an estimated Rabi frequency of 1 MHz, and the same 100 mW laser power as used experimentally. This gives a total off-resonance absorption of 89% of the pump light and absorption contrast of $C = 0.14\%$, in good agreement with our measurements.

FIG. 4. Experimental absorption contrast percentage as a function of microwave drive frequency and microwave power before the amplifier (Minicircuits ZHL-16W), measuring through the diamond with 100 mW of laser light ($I = 2.5 \times 10^4 \text{ W/m}^2$). The sample was used to test the absorption model using estimates of NV density and T_2^* from the observed linewidth (values in the main text). Note that due to input loss, we remain below the maximum gain threshold of the amplifier for all microwave powers shown, which is exceeded at +3 dBm.

B. Change in threshold current

We first calculate the lasing threshold current I_{th} with the diamond absent from the cavity. To do this, we fixed some of the parameters of the semiconductor gain medium. We choose a transparency carrier density of $N_{tr} = 1 \times 10^{25} \text{ m}^{-3}$ in the range typical for InGaN laser structures [52], a gain region volume of $V = 1.25 \times 10^{-16} \text{ m}^3$ ($25 \mu\text{m} \times 100 \text{ nm} \times 100 \mu\text{m}$), zero total cavity absorption $\alpha_t = 0$, a typical differential gain factor $a = 5 \times 10^{-20} \text{ m}^2$, a confinement factor Γ of 2%, and a carrier lifetime $\tau_N = 4 \text{ ns}$, giving a reasonable lasing threshold current of 50 mA [66]. We take the relation between the gain and the carrier density to be linear, with the carrier density close to transparency. We make the simplifying assumption that due to the low power, running close to lasing threshold we do not encounter gain compression effects, such that the factor $\epsilon \rightarrow 0$. We also initially make the simplifying assumption that the spontaneous emission rate is low, with $\beta \rightarrow 0$ (the importance of this second assumption will be tested in the final section of this work). These assumptions allow the threshold current I_{th} to be calculated easily from Eq. (8). We define $L_r = 10 \text{ mm}$, which is sufficient to include the diamond and any necessary optics in a practical implementation. We set mirror reflectivity $R_3 = 0.99$ and collect laser output from transmission through mirror R_1 . We calculate reflectivity R_2 from the Fresnel equations assuming an $\text{In}_x\text{Ga}_{1-x}\text{N}$ -air interface with refractive index $n \approx 2.6\text{--}2.9$ for $\text{In}_x\text{Ga}_{1-x}\text{N}$ [67].

We impose two feasibility limits on the threshold current I_{th} . The first is that it should not exceed 300 mA, based on the limits discussed in the technical documentation, in order to maintain thermal stability and for practical heat sinking for

FIG. 5. (a) Threshold current as a function of differential gain factor a and confinement factor Γ for diamond D3. (b) Change in threshold current due to the diamond absorption contrast $C = 0.02\%$ for diamond D3. Here, external cavity length L_r was 10 mm and output mirror reflectivity $R_1 = 0.9$. (Both the x and y axes use a $\log_{10}$ scale.)

a miniaturized diode or gain chip medium. The second is that the change in the threshold caused by the diamond absorption must exceed the shot noise of the drive current. From Fig. 5, we can see that these are mutually exclusive objectives. Setting an absorption contrast $C = 0.2\%$ (diamond D3) and output mirror reflectivity $R_1 = 0.9$ in order to achieve laser output while keeping the threshold current reasonably low, a low confinement factor Γ and high differential gain coefficient a result in the highest change in the threshold current ΔI_{th} and thus the strongest effect for sensing, but for very high I_{th} . Conversely, a higher value of Γ or lower a gives lower I_{th} , but ΔI_{th} shifts which are too small to be resolved.

Although N_{tr} is a factor usually defined by the semiconductor material, we note that the other parameters here which define I_{th} , Γ , a , and total cavity loss α_t including the mirror reflectivity and cavity output through R_1 , are all factors which are well understood and can be controlled and optimized at either the semiconductor growth stage or in the external cavity design.

C. Simulated ODMR

Here we calculate the ODMR spectrum that would be produced from the external cavity laser. We model a single microwave resonance from a single $m_s = 0 \rightarrow m_s = \pm 1$ transition using a Lorentzian line shape typical of ODMR for diamond [18]. We center our resonance at 2.82 GHz, replicating an ODMR resonance feature associated with a single NV axis, split from resonance features from other axes by an arbitrary weak DC offset magnetic field. The maximum amplitude is defined by the maximum change in threshold current between on and off microwave resonance and full width half maximum linewidth f_l . For simplicity, we assume that we can reach the pulsed readout linewidth defined by T_2^* . We calculate the external cavity laser output power using Eq. (10). Figure 6(a) shows the simulated ODMR for diamond D3, with a laser power output in the mW range for reasonable values of $\Gamma < 0.1$ and $a = 10^{-17}$ – 10^{-21} m². The equivalent plots for diamonds D1 and D2 are given in the Supplemental Material [34], with maximum power outputs in the range of nW and μ W, respectively. Unlike for conventional red fluorescence ODMR, the spectrum using this method is a peak at microwave resonance with zero background, rather than a small percentage change on a bright background.

D. Magnetic field sensitivity

We calculate the sensitivity to a magnetic field by taking the background noise level, dividing by the maximum ODMR slope, and by assuming a maximum frequency shift of ≈ 28 Hz–1 nT [13]. In our model, the primary sources of noise are readout from the photodetector and the noise on the drive current. The ultimate limit on both of these is shot noise of the output laser light and the drive current shot noise. We make no account for other direct sources of noise which are difficult to quantify, such as vibration or temperature fluctuations. Figure 7 shows a plot of the sensitivity for diamond D3 versus laser diode parameters for (a) the optical shot noise and (b) drive current shot noise limited regimes. We highlight that these calculated values neglect any effects from amplified spontaneous emission near threshold, which we explicitly include in Sec. III E. We note that in practice, the shot noise limited operation may be experimentally difficult to realize and we include an estimate based on a commercial current source with ppm-level noise in the Supplemental Material [34].

The noise limitations as a function of diode parameters are highlighted in Fig. 8 for diamond D3. Here, regions A and C represent where the operation is noise limited, and region B represents the region in which the system can operate. In region A, the change in the threshold current is less than the shot noise of the laser drive current ($\Delta I_{th} < I_{sh}$). In region C, the threshold current $I_{th} > 300$ mA exceeds a reasonable maximum drive current in order to maintain thermal stability. The limitations we impose mean that diamonds D1 or D2 have no viable operating region. For completeness, their sensitivity plots are included in the Supplemental Material [34].

By solving the rate model and calculating for laser diode output, we can calculate the sensitivity to a magnetic field for any valid physical parameters of the system, regardless of whether a diamond can be created with the requisite prop-

FIG. 6. (a) Simulated ODMR for diamond D3 at a range of differential gain factors $a = 10^{-21} \rightarrow 10^{-19.5}$ m² (exponents given in legend) measured by calculating the external cavity laser output power P as a function of microwave frequency for a Lorentzian line shape transition centered at 2.83 GHz and of linewidth defined by $f_l = \frac{1}{\pi T_2^*} = 3.2$ MHz. (b) Maximum laser power output P_{max} on resonance as a function of gain coefficient a for confinement factor $\Gamma = 0.01, 0.05, 0.1$. (c) The lasing threshold current for $I_{th} < 300$ mA, for the same three values of Γ . [The x axis in (b) and (c) uses a $\log_{10}$ scale.]

erties. This includes whether a value of T_2^* can be realized for a corresponding N_{NV} , making no assumption regarding the relation between these parameters or whether N_{NV} can be realized experimentally. Here we choose parameters R_1 , Γ , T_2^* , a , and NV⁻ density N_{NV} as the optimization variables, while fixing the diamond thickness ($d = 500$ μ m), Rabi frequency ($\Omega_R = 1$ MHz), laser beam width (0.5 mm), power (200 mW), and mirror reflectivities. We limit our laser power to 200 mW based on our rate model calculations to ensure that the majority of light is absorbed by the NV⁻ defects. We optimize using standard gradient descent methods. Figure 9 shows a plot of optical shot noise limited field sensitivity as a function of T_2^* and NV density. Sensitivity increased with higher T_2^* as would be expected, with maximum sensitivity at NV density of 10 ppm, above which high overall absorption by the diamond

FIG. 7. (a) Optical shot noise limited sensitivity for diamond D3 within a viable range for diode parameters a and Γ . Sensitivity is in the picotesla range, enabled by the elimination of the high noise from the background in the conventional fluorescence detection scheme. (b) The ultimate sensitivity limit imposed by shot noise on the laser drive current, worse by up to two orders of magnitude. These plots do not consider practical viability, with threshold currents > 4 A at low Γ . Zoomed-in plots in the low- Γ , low- a range are included in the Supplemental Material [34] for clarity. (Both the x and y axes use a $\log_{10}$ scale.)

acted to excessively reduce laser output. Subpicotesla level sensitivity is predicted for $T_2^* > 1 \mu\text{s}$ (0.3 – 0.02 pT/ $\sqrt{\text{Hz}}$ for $T_2^* = 1$ – $10 \mu\text{s}$). Here the optimal parameters were $a = 1.6 \times 10^{-20} \text{ m}^2$, $R_1 = 0.154$, and $\Gamma = 0.025$. These are parameters within the achievable range for a semiconductor gain medium (see Table I). We note that current demonstrations show that achieving long T_2^* times (e.g., $10 \mu\text{s}$) becomes impractical for high NV densities (e.g., 10 ppm), as detailed by Fig. 2, due to the impact of N - N interaction.

We note that in general, the highest sensitivity is realized for the lowest differential gain factor a . A standard laser diode demands a large a , maximizing gain vs carrier density (steeper output power vs drive current slope). Our scheme requires the reverse: that a small change in gain produced by the diamond on-off microwave resonance results in a large change in N_{ih} and I_{th} . In this respect, a quantum well structure with a flatter

FIG. 8. Regions where the sensor can and cannot operate due to imposed limitations. In regions A and C, operation is constrained by having a change in threshold that is less than the shot noise of the drive current and $I_{th} > 300 \text{ mA}$, respectively. In region B, operation is possible. (Both the x and y axes use a $\log_{10}$ scale.)

logarithmic relation between gain and carrier density would seem preferable. However, as we demonstrate in Fig. 10, using our model, with modifications to the phenomenological

FIG. 9. Best field sensitivity optimizing variables listed in the main text as a function of (a) NV^- density and (b) T_2^* in μs , with the inset showing a zoomed plot at the highest simulated values of T_2^* . The best sensitivity was observed at the highest T_2^* , for NV^- density of 10 ppm . Above this, the total absorption for the diamond was too high, limiting laser output and sensitivity. [The x and y axes in (a) and x axis in (b) use a $\log_{10}$ scale.]

FIG. 10. Calculated magnetic field sensitivity (left y axis) and laser diode threshold current (right y axis, dashed) as a function of differential gain factor a using an empirical model for a quantum well laser diode. The lasing threshold current increases such that subpicotesla sensitivity is not reached at a feasible threshold current (< 300 mA). Here we take confinement factor $\Gamma = 0.01$ as an example of the low values that are typical of a quantum well laser diode. (The x and y axes use a $\log_{10}$ scale.)

description of the medium gain (detailed in the Supplemental Material [34]) and with the same optimization methodology as above results in the threshold current exponentially exceeding drive current feasibility limits before subpicotesla/ $\sqrt{\text{Hz}}$ sensitivity is reached, for any typical value for Γ in the low percentage range.

E. Effect of spontaneous emission

In the previous sections and past literature, the physical role of spontaneous emission in the semiconductor gain medium that was used was not considered. In order to maximize sensitivity, it is necessary to operate at or close to the off-resonance lasing threshold. Without spontaneous emission, this can be treated as a step cut-on, with zero or near-zero emission before lasing begins at I_{th} . With spontaneous emission included, modeled by finite β in Eqs. (4) and (5) above, the power-current relationship close to threshold instead follows a shallow curve, resulting from weak amplification of spontaneous emission near threshold producing light emission below I_{th} . Figure 11(a) shows this effect for varying β . This acts to severely limit sensitivity [Fig. 11(b)] by reducing the contrast and adding background shot noise. Typical values of β range from 10^{-3} to 10^{-5} , depending on the laser diode structure. We estimate approximately an order of magnitude worse sensitivity at the low end of this range than with $\beta = 0$.

IV. CONCLUSION

In this work, we propose a scheme for laser threshold sensing using an external cavity laser configuration with a current driven semiconductor lasing medium. Using the change in lasing threshold, light emission only occurs on microwave resonance. This eliminates the bright background that limits sensitivity using conventional red fluorescence emission. Predicted sensitivities (summarized in Table II) for magnetometry with realistic cavity parameters and intrinsic material parameters are in the $\text{pT}/\sqrt{\text{Hz}}$ range, offering a route to

FIG. 11. (a) External cavity laser output P as a function of semiconductor laser medium drive current I , varying spontaneous emission factor β . The result of increasing β is that there is no longer a sharp lasing cut-on at the threshold. (b) The effect on the achievable sensitivity of this effect, with sensitivity considerably reduced by up to two orders of magnitude for $\beta = 10^{-2}$.

improvement over existing methods. We find that these sensitivities may be reachable at far lower laser power (< 200 mW) than realized in equivalent fluorescence-based diamond bulk sensing schemes, including our recent experimental work which utilized up to 2 W of green pump laser to obtain equivalent sensitivities in the tens of pT range [68]. This significant decrease of laser power may facilitate sensor miniaturization and thus the implementation of portable sensing schemes.

Our model has limitations: we base our calculations on emission into a single laser mode and do not calculate the dynamics of the system, such as rapid switching in a pulsed operation scheme. Although beyond the focus of this work, we note that the latter may be a promising route for future investigation. A scheme where the laser medium could be initially pumped and then the pump shut off while retaining population inversion during sensing, typical of a Q-switched setup, would only be limited by the optical shot noise of any emitted laser light. This is, however, challenging to achieve for a semiconductor laser due to the short excited state (carrier) lifetime.

A key physical limitation of any laser threshold scheme is the role of amplified spontaneous emission. This blurs the sharp lasing transition, giving nonzero light emission even below threshold and compromising sensitivity. A broad transition can be avoided by minimizing gain factor β , although

TABLE II. Summary of key values from our calculations for modeled diamonds D1–D3 and an optimal (ideal) diamond, including the assumed NV density, T_2^* , maximum shift in threshold current, and best calculated sensitivities within the modeled parameter ranges of a , Γ , with and without spontaneous emission (SE) taken into account. We present the best sensitivity within the modeled range limited by either the optical shot noise (SN) of the laser emission or limited by the shot noise of the drive current (brackets). For D1 and D2, * denotes not viable, as the drive current shot noise exceeds the change in threshold ΔI_{th} .

Diamond	NV density (ppm)	T_2^* (μ s)	Maximum threshold shift (μ A)	Best sensitivity, without SE, optical (current) SN limit pT/ $\sqrt{\text{Hz}}$	Best sensitivity, including SE, optical SN limit, $\beta = 10^{-5} \rightarrow 10^{-3}$ pT/ $\sqrt{\text{Hz}}$
D1	10^{-3}	5	0.00122	4.6 (1834*)	
D2	0.1	0.75	41.4	3.1 (124*)	
D3	10	0.1	6.2×10^3	4.3 (53)	22–73
Ideal	10	1–10	7.3×10^3 – 8.4×10^3	0.3–0.02 (-)	

this is difficult for a semiconductor laser, particularly since β can scale inversely with the size of the gain medium [69]. Obtaining a gain chip or antireflective coated laser diode with the right parameters is challenging, especially for green wavelengths. This problem also exists for infrared absorption since the laser emission must match the 1042 nm gap in the singlet state. In our scheme, running in the infrared could be achieved by extending the external cavity design proposed here using a diffraction grating in a Littrow and Littman-Metcalf configuration to create a tunable system.

We consider, in this work, a normal incidence beam path through a fixed diamond thickness $d = 500 \mu\text{m}$. We note that a higher sensitivity within feasible limits of threshold current could potentially be reached with a thinner diamond with a very high NV density. However, in the limit of $d \rightarrow 0$, other effects not considered in our model may act to limit performance, such as variation in NV density and the role of other types of defects. Measurements of absorption and T_2^* as a function of diamond thickness would be extremely useful in determining behavior in this regime. We also consider that it may be possible to reach higher sensitivity in the low NV density regime using an extended beam path achieved through internal reflection in a thicker diamond. This again requires new experimental measurements to precisely quantify the losses (due to reflection or absorption) in such a geometry.

We note that the fundamental limit for the scheme is the level of contrast C generated between the on-off microwave resonance states, which is very low for a large diamond ensemble. However, the scheme is not specifically limited to diamond and is broadly applicable for any material where a large enough, controllable difference in optical absorption could be generated. The advantage of using diamond is the ability to coherently manipulate the desired states in a quantum sensing scheme. Our calculations indicate the scheme will likely only work for diamonds with a high (> 1 ppm) NV⁻ density. Such diamonds have a worse ensemble T_2^* time, limited by nitrogen spin interaction. A developing solution here may be to use optimal control methods in order to better control the ensemble. Such methods are widely implemented for nuclear magnetic resonance and electron spin resonance on bulk samples, but have yet to be fully developed for sensing using diamond defects [70].

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