

Low-Charge-Noise Nitrogen-Vacancy Centers in Diamond Created Using Laser Writing with a Solid-Immersion Lens

Viktoria Yurgens,* Josh A. Zuber, Sigurd Flågan, Marta De Luca, Brendan J. Shields, Ilaria Zardo, Patrick Maletinsky, Richard J. Warburton, and Tomasz Jakubczyk



Cite This: *ACS Photonics* 2021, 8, 1726–1734



Read Online

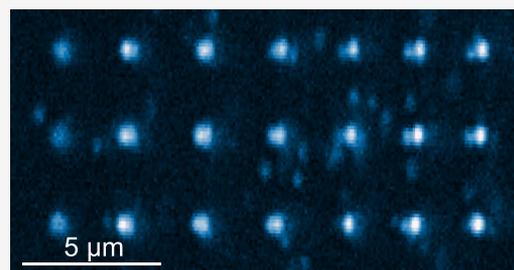
ACCESS |

Metrics & More

Article Recommendations

ABSTRACT: We report on pulsed-laser-induced generation of nitrogen-vacancy (NV) centers in diamond facilitated by a solid-immersion lens (SIL). The SIL enables laser writing at energies as low as 5.8 nJ per pulse and allows vacancies to be formed close to a diamond surface without inducing surface graphitization. We operate in the previously unexplored regime, where lattice vacancies are created following tunneling breakdown rather than multiphoton ionization. We present three samples in which NV center arrays were laser-written at distances between ~ 1 and $40 \mu\text{m}$ from a diamond surface, all presenting narrow distributions of optical linewidths with means between 62.1 and 74.5 MHz. The linewidths include the effect of long-term spectral diffusion induced by a 532 nm repump laser for charge-state stabilization, thereby emphasizing the particularly low-charge-noise environment of the created color centers. Such high-quality NV centers are excellent candidates for practical applications employing two-photon quantum interference with separate NV centers. Finally, we propose a model for disentangling power broadening from inhomogeneous broadening in the NV center optical linewidth.

KEYWORDS: diamond, nitrogen-vacancy center, laser writing, solid-immersion lens, quantum information processing



The negatively charged nitrogen-vacancy (NV) center in diamond is among the most promising solid-state system implementations of a quantum bit.^{1,2} Its applications include sensing^{3,4} and quantum communication, with progress demonstrated in spin-photon⁵ and long-distance spin–spin entanglement.⁶ However, interconnecting many NV centers for large-scale quantum networks suffers from the low generation rate of indistinguishable photons from individual NV centers.⁶

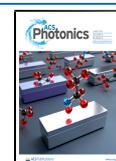
Fabrication of any diamond-based photonic device requires structuring of the diamond on a submicron scale. An example is the open Fabry–Perot microcavity—a platform enabling easy optical access, mode-matching, and *in situ* tuning of the cavity resonance to an emitter.^{7–10} In this case, bulk diamond is thinned down to a few-micron-thick membrane, a minimally invasive fabrication.^{11–16} The technique is a promising route to enhance radically the generation rate of indistinguishable photons from individual NV centers. However, even such minimal processing has so far resulted in degradation of the NV centers' optical quality, manifesting in a large spectral diffusion.^{13,17} The random spectral fluctuations in the zero-phonon line (ZPL) frequency are caused by charge noise present in the structured crystal: NV centers exhibit a large change in the static dipole moment between the ground and excited state,¹⁸ making the ZPL emission frequency strongly dependent on the local electric field.¹⁹ The electric field in turn depends on the spatial configuration and occupation of the surrounding charge

traps in the form of defects and impurities. Methods of NV center creation that do not result in the formation of parasitic defects and impurities are therefore desired.

The creation of an NV center typically involves displacing a carbon atom from its lattice site to create a vacancy. This occurs above a threshold energy of 35 eV.²⁰ While proximity to the surface is required for photonic applications, it is at the same time desirable to form the NV centers at a depth of at least tens of nanometers from the diamond surface in order to minimize the impact of the surface charge and magnetic noise on the NV centers' spin and optical coherence. The challenge is therefore to provide this energy inside the crystal lattice. Electrons or ions with kinetic energies far above keVs readily provide such an energy to depths from a few nanometers to tens of micrometers and above in the case of electrons.²¹ However, most of this energy is released via collisions on the particle's trajectory, leaving extended damage and presumably hard-to-anneal vacancy complexes. A widely employed technique to create NV centers at precisely controlled depths consists of nitrogen-

Received: February 19, 2021

Published: June 3, 2021



ion implantation followed by high-temperature annealing.²² The technique is well-suited for creating NV centers for sensing applications^{23,24} but has recently been shown to create NV populations where approximately half of the NV centers show spectral diffusion above a few GHz.^{25,26} In addition, the NV centers formed from the implanted nitrogen ions have on average much poorer optical quality than their counterparts formed from the native nitrogen,^{25,26} suggesting that vicinity to the fabrication-related damage plays a crucial role for the spectral diffusion.

Typically, the inhomogeneously broadened linewidth distribution in unprocessed bulk samples irradiated by ions or electrons is characterized by linewidths on the order of 100 MHz,^{17,25,26} disregarding half of the population with above-GHz linewidths for the nitrogen-implanted samples. After the samples are thinned down to less than a micron by reactive ion etching, most of the observed NVs are characterized by >1 GHz linewidths.¹³ This increase in linewidth is detrimental for photon indistinguishability.^{27,28} Good two-photon interference is only possible when the inhomogeneous broadening is small compared to the homogeneous linewidth. For an NV center in bulk diamond, the lifetime limit sets the smallest possible homogeneous broadening of just ~13 MHz,²⁹ far below typical inhomogeneous linewidths. However, in a resonant microcavity, the Purcell effect strongly reduces the lifetime, thereby increasing the lifetime limit. Nevertheless, in current devices, the inhomogeneous broadening is still higher than the Purcell-enhanced homogeneous linewidth.¹³ Radical progress in improving the emitter properties or microcavity properties, ideally both, is therefore required.

A recent study demonstrated progress in obtaining NV centers with linewidths of approximately 200 MHz in 3.8- μm -thick diamond membranes.¹⁷ In even thinner structures, linewidths down to 250 MHz were reported with a new approach based on implanting nitrogen after microstructuring²⁶ and also in samples structured with a slow etching procedure.³⁰ To the best of our knowledge, narrower linewidths than these have never been reported in micron-thick diamond. The origin of the decrease in the NV centers' optical quality is potentially related to the damage caused to the crystal lattice during implantation and etching, but the microscopies are not well-understood. The lack of optically coherent NV centers in structured diamond clearly points to the need for improved NV creation methods.

Laser writing has emerged as a promising technique for creating deterministically positioned NV centers in diamond³¹ and other color centers in various wide-bandgap materials.^{32–36} NV centers created through laser writing in ultrapure diamond are characterized by long spin coherence times, low spectral noise, and high charge stability.³¹ Crucially, their fabrication induces minimum damage to the diamond lattice, potentially reducing the charge noise in thin diamond membranes. These NV centers are formed from native nitrogen atoms, and implantation is avoided completely, meaning that the lattice is damaged only locally in the laser focus.

Initial studies demonstrated laser writing of NV centers using an oil-immersion lens in combination with wavefront correction.^{31,37} Here, we report on a different approach employing a standard air objective together with a solid-immersion lens (SIL), giving a high numerical aperture (NA) together with minimized aberrations, yielding a diffraction-limited laser focal volume inside diamond. This technique not only enables creation of NV centers with low pulse energies but

also makes it possible to laser-write in close proximity to the diamond surface, owing to the decreased laser intensity at the diamond–air interface. We demonstrate the high quality of the NV centers created with this method, with a record-low noise measured in the presence of a charge-state reinitialization pulse, and determine separately the inhomogeneous and power-broadening contributions to the ZPL linewidths.

LASER WRITING OF NV CENTERS

Laser writing of NV centers is based on irradiation of a diamond sample with single high-energy femtosecond pulses. The laser pulses create vacancies; NV centers form in a subsequent thermal annealing step, in which the mobile vacancies combine with nitrogen impurities in the diamond. Photoinduced damage in a transparent wide-bandgap material such as diamond requires the generation of excited free electrons through nonlinear mechanisms. This optical breakdown can result from tunneling or multiphoton absorption followed by avalanche ionization in the volume of the tightly focused laser beam.^{38–40} These processes displace single atoms locally and can create vacancies. In diamond, vacancy creation can be identified by the vacancy's spectral signature GR1.⁴¹ The dominant mechanism for the creation of the energetic seed electrons responsible for the optical breakdown can be determined using the Keldysh parameter⁴²

$$\gamma = \frac{\omega}{e} \sqrt{\frac{m_{\text{eff}} c n \epsilon_0 E_g}{I}} \quad (1)$$

where ω is the angular frequency of the laser, I is the laser intensity, m_{eff} is the electron effective mass, c is the speed of light in vacuum, n is the linear refractive index of the material, ϵ_0 is the vacuum permittivity, and E_g is the direct bandgap of the material. For $\gamma < 1$, tunneling breakdown is the primary process behind the creation of seed electrons, while for $\gamma > 1$, multiphoton ionization dominates. In the multiphoton ionization regime, owing to the strong nonlinearity of the process,³¹ the volume in which the atom displacement takes place can be much smaller than the size of the focal spot. However, during the annealing step, the created vacancies diffuse, and this process limits the NV placement accuracy, offering a precision down to hundreds of nanometers in initial reports.^{31,37}

To minimize the impact of the laser writing process on the crystal quality, the volume of the laser focus needs to be reduced by maximizing the NA of the focusing optics. However, spherical aberration induced at the air–diamond interface (with refractive indices of $n = 1$ and $n = 2.42$, respectively) also increases with increasing NA. We resolve this dilemma by using a truncated hemispherical cubic zirconia SIL (t-SIL) with a refractive index of $n = 2.14$, which both maximizes the NA and reduces the spherical aberrations (Figure 1(a,b)).

An absolutely crucial problem is that the energy of the ultrashort laser pulses required to create vacancies in bulk diamond typically exceeds the threshold at which the diamond surface is degraded.^{31,43,44} Creating vacancies in bulk diamond with laser writing is therefore only possible if the laser focus is spread over a significant area at the surface. To investigate this, we consider the case of imaging a laser with a wavelength of 800 nm with an objective lens of NA = 0.6 in Figure 1. In air, the extent of the focal spot along the optical axis, Δz , is in the best case limited by diffraction to a value of approximately 3.9 μm . This extent will be preserved roughly on focusing a few microns below the diamond surface. The large Δz is problematic: the

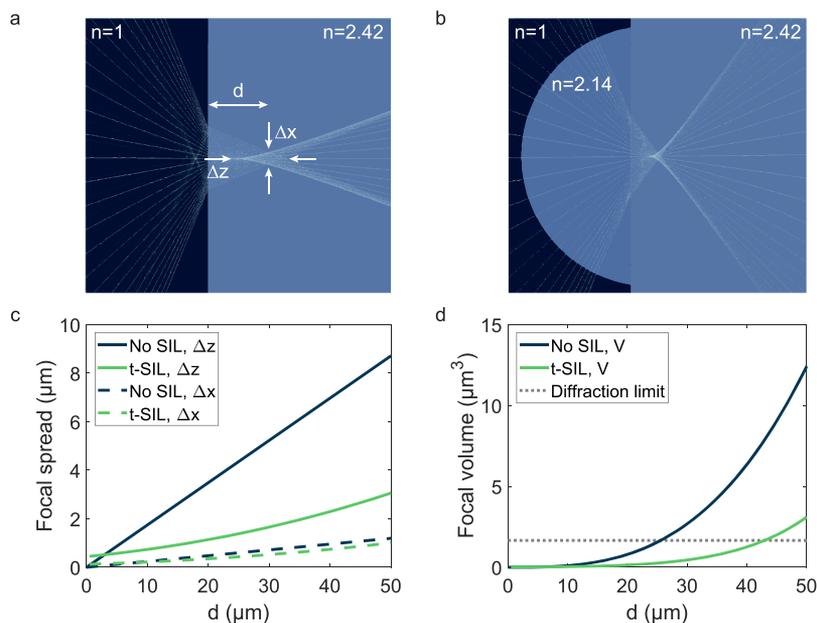


Figure 1. Geometrical optics simulation of the spread of rays on focusing through an air–diamond interface. (a) Spherical aberration for rays passing from air (left) to diamond (right) without corrective optics. Δx is the in-plane focal spread, and Δz is the focal spread along the optical axis; d is the distance from the point of minimum Δx to the diamond surface. (b) Spherical aberration for rays passing from air to diamond through a truncated hemispherical solid-immersion lens (t-SIL). (a,b) were calculated using the tool from ref 48. (c) Calculated focal spread in-plane (solid lines) and along the optical axis (dashed lines) for the cases without a SIL and with a t-SIL on passing through an air–diamond interface, as a function of the focusing depth d . (d) Calculated focal volume versus d without and with a t-SIL. (c,d) take a t-SIL truncation of 30 μm and an NA of 0.6. The geometrical optics simulations ignore diffraction, which determines the focal volume when aberrations are absent. In (d), the diffraction limit is estimated as $(\Delta x)^2 \Delta z$ with $\Delta x = 1.2\lambda/(n\text{NA})$ and $\Delta z = 2.0\lambda/(n\text{NA}^2)$, with $n = 2.42$.

intensity at the surface is large, and as a result, surface graphitization will occur before vacancy creation within the diamond. Focusing at a larger depth does not solve this problem. Geometrical optics shows that the spherical aberrations cause Δz to increase with depth d (Figure 1(c))—the focal spot becomes so elongated along z that surface graphitization is likely to remain a problem. This corresponds to our experience—without a t-SIL, surface graphitization occurred before vacancy creation. In the limit that the t-SIL is made of diamond, spherical-aberration-free imaging is achieved at a depth d , which matches the truncation t . Furthermore, the diffraction limit reduces simply because the relevant wavelength is the wavelength in the diamond, λ/n . For NA = 0.6, Δz reduces to 1.6 μm . These considerations suggest that vacancy creation is possible at depths starting at about 2 μm from the diamond surface, and it motivates the use of the t-SIL. In practice, the t-SIL has a slightly lower refractive index than diamond (such that a spherical aberration is introduced at the t-SIL–diamond interface), and it is impractical to use a different t-SIL for each depth. We find however that a single t-SIL gives good imaging out to a depth up to 50 μm (Figure 1(c,d)). The main improvement over imaging without the t-SIL is the reduction in Δz , crucial in order to keep the intensity at the surface small.

An additional advantage is that the t-SIL improves the photon collection efficiency and thereby the sensitivity to the weak signals emitted by the created vacancies by reducing the number of photons lost due to total internal reflection at the diamond–air interface.^{45–47} Moreover, the implementation of a t-SIL is both cost-effective and easily implemented. A limitation is perhaps the writing area—excellent imaging is achieved only close to the center of the t-SIL. However, a writing area of approximately 25-by-25 μm^2 is available for a t-SIL with a diameter of 0.5 mm, sufficiently large for many purposes.

We perform laser writing in a room-temperature home-built confocal microscope with a separate injection path for the femtosecond pulsed laser. A 532 nm Nd:YAG laser is used for nonresonant excitation of the NV centers. Photoluminescence (PL) is filtered by a long-pass filter (Semrock, 594 nm RazorEdge) and collected by an avalanche photodiode (APD) (Excelitas, SPCM-AQRH-15-FC) or a liquid-nitrogen-cooled charge-coupled device (CCD) camera coupled to a grating spectrometer (Princeton Instruments). We use a Spectra Physics Spirit 1030-70 fs ytterbium-doped fiber laser together with a Spirit-NOPA-2H noncollinear optical parametric amplifier, which together create pulses at a wavelength of 800 nm with a duration of approximately 35 fs. A half-wave plate and a Brewster-angle polarizer are used for tuning the pulse energies. The flat side of a t-SIL (radius 500 μm , truncation between 0 and 50 μm) is polished using a polishing suspension containing 50-nm-sized alumina particles. The t-SILs are then placed on the diamond samples together with an optical coupling gel in an attempt to reduce aberrations resulting from interface imperfections. We focus the 800 nm laser with a standard air objective (Olympus, MPLFLN100x, NA = 0.9 or Olympus, LCPLFLN100xLCD, NA = 0.85). The resultant NA including the effect of the t-SIL is about 1.8, slightly higher than that used in ref 31. We note however that the full-width-at-half-maximum (fwhm) of the beam is about the same as the input aperture of the objective, reducing the contribution of the rays at the highest angle. The performance of the optical setup including the t-SIL was determined by confocal imaging of the emission from single, pre-existing, shallow NV centers. The fwhm of a single bright spot was measured at 125 ± 3 nm. This value is just above the theoretical minimum of 121.1 nm, as given by $\frac{0.52}{\text{NA}} \frac{\lambda_1 \lambda_2}{\sqrt{\lambda_1^2 + \lambda_2^2}}$ for the confocal configuration (using $\lambda_1 = 532$ nm and $\lambda_2 = 700$ nm

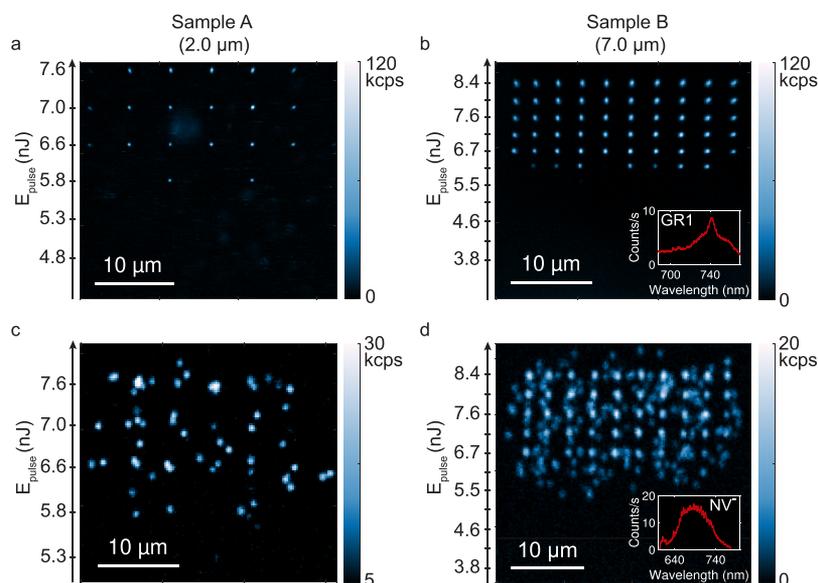


Figure 2. (a,b) Confocal scans of samples A and B (indicating their respective NV creation depths and pulse energies) showing GR1 photoluminescence (inset in b) at room temperature before annealing. The scans were performed through a t-SIL. (c,d) Confocal scans of samples A and B, showing NV⁻ photoluminescence (inset in d) at room temperature after annealing to 1100 °C. The scans were performed without a t-SIL.

for the excitation and main NV phonon-sideband emission wavelength, respectively, and taking $NA = 0.85 \cdot 2.14$). This close agreement between the measured spot size and the diffraction limit demonstrates that the optical components, notably the objective and the t-SIL, do not introduce unwanted aberrations.

We follow the experimental procedure described in ref 31 to create arrays of vacancies in bulk electronic-grade diamond samples (40 μm thick, $[N] < 5$ ppb, Element Six). We use single pulses at increasing pulse energies (3.8–35.8 nJ) and subsequently anneal the diamond samples at 1100 °C to create NV centers. Surface markers for locating the laser-written arrays after annealing are made through graphitization of the diamond surface by increasing the pulsing frequency of the femtosecond laser to 100 kHz, removing the t-SIL, and increasing the energy to a value above the graphitization threshold of the surface (typically around 5 nJ).

Results on two diamond samples are shown in Figure 2, with arrays of vacancies (a,b) and NV centers (c,d), respectively, created with increasing pulse energies from the bottom of the arrays. The scans were recorded at room temperature. The arrays were made at depths of 2.0 and 7.0 μm below the diamond surface, respectively, where we note that the former demonstrates that our method can be used to create NV centers in close proximity to the diamond surface without inducing graphitization. Vacancy creation at shallower depths was not possible without damaging the diamond surface. This observation aligns with the analysis of the dimensions of the focal spot. For sample A, pulse energies of between 4.8 and 7.6 nJ were used, with visible GR1 photoluminescence (inset, Figure 2(b), 100 s integration time) appearing for an energy of 5.8 nJ per pulse; for sample B, pulse energies between 3.8 and 8.4 nJ were used, with visible GR1 photoluminescence appearing for an energy of 6.1 nJ per pulse. All pulse energies refer to values at the output of the microscope objective. Both samples were annealed in vacuum for 3 h at 1100 °C, after which the spectral signatures of negatively charged NV centers (inset, Figure 2(d), 100 s integration time) were observed.

Calculation of the Keldysh parameter γ for these vacancy-creation threshold energies, with the specified pulse duration and focal volume, gives $\gamma \ll 1$, meaning that we operate in a regime where tunneling breakdown is dominant over multi-photon ionization, in contrast to previous work.³¹ Tuning of the vacancy-creation regime could be carried out by changing the laser writing wavelength and increasing the pulse length, but with the latter shrinking the window for NV center creation due to a decrease of the graphitization threshold.⁴⁹

Sample B shows a clear array-like NV pattern after annealing, where the main array points contain multiple NV centers, as demonstrated by the photon count in the confocal scans and by the existence of more than two lines in subsequent photoluminescence excitation (PLE) measurements. The diffusion of vacancies during annealing is also clearly visible—single NV centers form up to several hundreds of nanometers from the array spots, similar to the results of previous experiments.^{31,50,51} Lower pulse energies and a larger spatial separation between the focal spots were therefore used for sample A in order to obtain predominantly spots with single rather than multiple NV centers.

Vacancy photoluminescence after laser writing was observed both with the t-SIL and after displacing the t-SIL from the laser-processed area. If no t-SIL was used during the laser-writing process, neither graphitization nor GR1 photoluminescence could be detected in the bulk of the sample up to a pulse energy of 52 nJ (highest available in our experimental configuration) and a continuous exposure with a pulse repetition rate of 1 MHz with durations of up to tens of seconds. Similar observations were made after exposures performed with a t-SIL but without using the index-matching gel. Also after annealing, there was no NV photoluminescence in areas exposed without a t-SIL. These observations indicate that the use of a t-SIL dramatically reduces the threshold for vacancy generation and can act as the key element enabling laser-induced NV formation using fs laser sources.

A third sample, sample C, was patterned in a way to create NV centers as close to the diamond surface as possible without inducing visible damage.⁵² PL images of the resulting array of

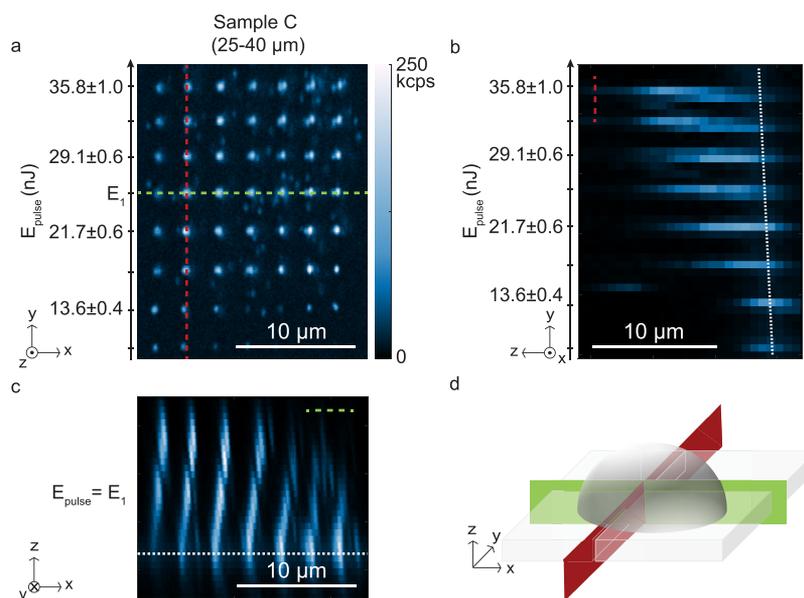


Figure 3. (a) Confocal scan of NV centers in sample C, laser-written in an inverse geometry where laser pulses were focused close to the bottom surface of the diamond. Each consecutive spot along a line was laser-written at a reduced distance from the surface compared to its neighbor. The color bar applies to a–c. Some of the spots contain multiple NV centers—this accounts for the changes in brightness from spot to spot. (b) Cross section in the plane indicated by the red dashed line in (a). The white dashed line indicates the bottom diamond surface, determined by the position at maximum intensity of the reflected excitation laser. (c) Cross section in the plane indicated by the green dashed line in (a), for a pulse energy $E_1 = 25.7 \pm 0.5$ nJ. The white dashed line indicates the bottom diamond surface. (d) Schematic of the two cross sections through the diamond, with a t-SIL placed on top.

NV centers are shown in Figure 3(a–c). The increased energy density on the top diamond surface upon focusing the laser less than $\sim 2 \mu\text{m}$ from it typically resulted in graphitization of the diamond surface or damage of the t-SIL. For this reason, an inverse geometry was chosen. In this geometry, the laser pulses were applied through the t-SIL and diamond but focused close to the diamond back surface, the one further from the t-SIL. Several NV arrays were patterned at different depths in the diamond in this configuration. An objective lens with $\text{NA} = 0.85$ and a correction ring were used for this study to further minimize the spherical aberration; the correction ring was set to maximize the laser focus intensity at the bottom surface of the sample.

NV centers could be created in the full range of pulse energies in Figure 3(a), from 7.6 ± 0.4 to 35.8 ± 1.0 nJ. Even at the highest pulse energy, there were no signs of graphitization. This energy window is remarkably wide compared to previous studies.^{31,37} We propose that this is a key consequence and advantage of NV center laser writing in the tunneling regime instead of the multiphoton ionization regime. In order to test NV writing at various depths and in order to find a possible distance limit from the surface for NV creation, each array additionally included a smaller depth variation of $0.4 \mu\text{m}$ between each laser writing spot within a row. This resulted in NV centers covering distances from ~ 1 up to $15 \mu\text{m}$ from the bottom diamond surface, corresponding to laser writing depths of between 25 and $40 \mu\text{m}$ from the top surface. Figure 3(b,c) shows cross sections through the diamond to illustrate the resulting slightly tilted layers. In Figure 3(b), it is apparent that a lower pulse energy was needed to create the NV centers closest to the surface, as compared to writing NV centers further inside the bulk. The reason for this is presently unknown. Figure 3(d) illustrates the cross sections and the general t-SIL-on-diamond geometry.

■ LINEWIDTH CHARACTERIZATION

We characterize the ZPL linewidths through PLE scans in a liquid helium bath cryostat. The NV centers are resonantly excited using a tunable narrow-linewidth external cavity diode laser (New Focus Velocity TLB-6704) using powers between 15 and 450 nJ and the linewidths are determined by sweeping the excitation frequency over the ZPL resonance while recording the photons emitted into the phonon sideband. Each line is measured at the lowest possible power that gives a good signal-to-noise ratio, in order to have as little power broadening as possible. Between the periods of excitation and readout, a negative charge-state repump is carried out with a 532 nm Nd:YAG-laser with an average power of 0.6 mW. The full cycle of repump followed by resonant excitation and readout is repeated at a frequency of 100 kHz, with a 21-to-68 green-to-red laser duty cycle. The inset in Figure 4(a) shows the PLE measurement sequence.

Two PLE scans are presented in Figure 4(a,b), showing ZPL resonances with Lorentzian lineshapes with fwhm linewidths of 35.9 ± 0.9 , 34.5 ± 1.8 , and 32.6 ± 2.3 MHz, close to the lifetime-limited linewidth of 13 MHz measured for a native NV center in a bulk natural diamond sample.²⁹ (In the low-power regime, a Gaussian fits the PLE data only slightly better than a Lorentzian: average mean-standard-errors of 0.014 and 0.017, respectively. We use here a Lorentzian for simplicity.) We note that ref 29 measured linewidths between the repump pulses and thereby excluded a major source of inhomogeneous broadening, charge reconfiguration of the environment on resetting the NV center charge. In our case, we integrate over many repump cycles, thereby including all the contributions to the inhomogeneous broadening. We note also that fitting the lines measured at low power (where inhomogeneous broadening dominates) with Gaussian functions rather than Lorentzians yielded an average difference in fwhms of only 1.5% for the three samples, further justifying the use of a Lorentzian function to describe the

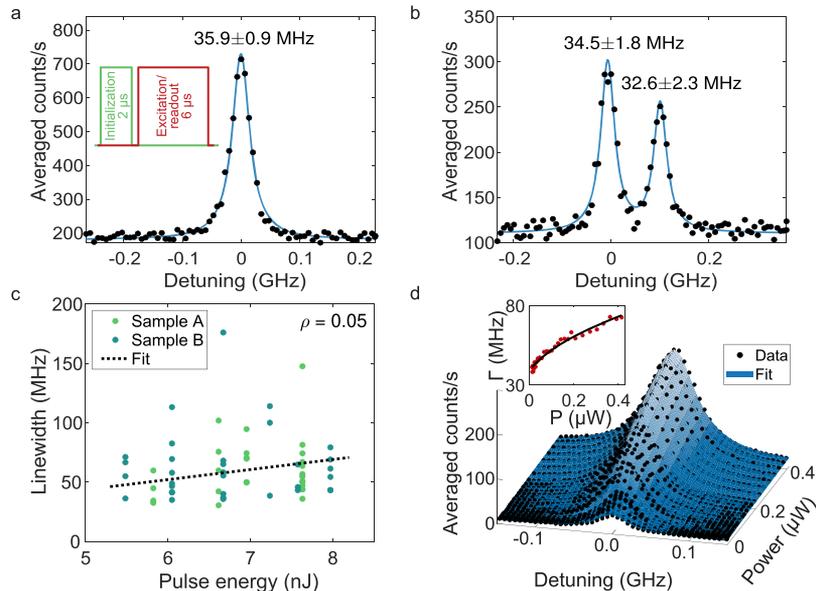


Figure 4. (a) Optical linewidth measured on an NV in sample A. Zero detuning corresponds to a ZPL frequency of 470.494 THz. The inset shows the measurement sequence for the photoluminescence excitation measurement. (b) Linewidth measured on an NV center in sample B. Zero detuning corresponds to a ZPL frequency of 470.509 THz. (c) Dependence of the linewidths in the two samples on the fs laser pulse energy used in the laser writing process. Each data point corresponds to a separate NV center, except for a few cases where two lines were measured for one NV center. The dotted line is a linear fit with a Pearson's $\rho = 0.05$, indicating little to no correlation between linewidth and pulse energy. (d) Resonant power dependence of the ZPL of an NV in sample A, fitted according to eq 3. The inset shows the dependence of the extracted fwhm linewidth on power, fitted following eq 4 and demonstrating clear power broadening.

inhomogeneous broadening. Figure 4(c) shows the dependence of the linewidths on the femtosecond laser pulse energy, where the dotted line is a linear fit to the data. A Pearson's $\rho = 0.05$ for the fit indicates no significant correlation between ZPL linewidth and pulse energy, in contrast to ref 31.

Figure 4(b) shows a splitting between the resonances of just 109 MHz, which indicates a low strain level in the surroundings of the specific NV. (Here, we make the reasonable assumption that the peaks correspond to the E_x and E_y transitions of the same color center.) In general, the three laser-written samples demonstrate several doublets with exceptionally low peak splittings down to 81 MHz, but a more systematic study is required in order to obtain a definite conclusion on the strain levels of laser-written NV centers and to exclude the possibility that the two lines originate from separate NVs (however, such a coincidence in spatial location, linewidth, and ZPL energy between different NVs is highly unlikely). The absolute ZPL frequency has a distribution centered at 470.49 THz with a standard deviation of 14 GHz, similar to the distribution in ref 25.

To disentangle inhomogeneous broadening of the linewidths (Γ_{in}) from power broadening characterized by the Rabi coupling Ω , we perform a systematic study of the linewidth as a function of resonant power. To simplify the analysis, we assume a Lorentzian spectral diffusion shape characterized by fwhm Γ_{in} : the probability of the emitter frequency being equal to f^* is given by the (normalized) Lorentzian function $L(f^* - f_0, \Gamma_{\text{in}})$ where f_0 is the average emitter frequency. The occupation of the excited level of a driven two-level system with radiative decay rate γ is given by⁵³

$$\rho_{22} = \frac{\left(\frac{1}{2}\Omega\right)^2}{4\pi^2(f - f^*)^2 + \left(\frac{1}{2}\gamma\right)^2 + \frac{1}{2}\Omega^2} \quad (2)$$

where $\Omega = \sqrt{c \cdot P}$, with P being the excitation power and c being an effective coupling strength that depends on the laser's focal volume, the laser incoupling efficiency, and the NV center's dipole moment and orientation. The experimentally measured lineshapes can be described by a convolution of L with ρ_{22} , yielding an expression for the counts as a function of frequency in a PLE measurement

$$C(f) = \frac{A}{4\pi} \frac{\Omega^2}{\sqrt{\gamma^2 + 2\Omega^2}} \frac{\frac{1}{2}\Gamma}{(f - f_0)^2 + \left(\frac{1}{2}\Gamma\right)^2} \quad (3)$$

where A depends on the setup's collection efficiency and the average NV photon emission rate, the "dead" time during the repump pulse, and the time spent in the NV^0 charge state as well as the time spent in spin states that are not cycled with the resonant laser.⁵⁴ The fwhm linewidth is

$$\Gamma = \Gamma_{\text{in}} + \frac{\sqrt{\gamma^2 + 2\Omega^2}}{2\pi} \quad (4)$$

where we assume $\gamma = 13$ MHz.

Figure 4(d) shows a PLE measurement as a function of the resonant excitation power, and the data can be fitted well with eq 3. Power broadening is clearly visible. The inset shows the extracted fwhm linewidth as a function of power, fitted with the linewidth described by eq 4. The full model gives a value of $c = (1.0 \pm 0.1) \cdot 10^5 \text{ MHz}^2/\mu\text{W}$ for the effective coupling strength and a value of $\Gamma_{\text{in}} = 25.5 \pm 1.2$ MHz for the inhomogeneous broadening.

Figure 5 shows a summary of the linewidth measurements on samples A, B, and C. All three samples demonstrate narrow distributions of linewidths, with mean linewidths (m) of 62.1, 62.1, and 74.5 MHz and standard deviations (SD) of 23.8, 24.6, and 50.3 MHz, respectively, when assuming log-normal

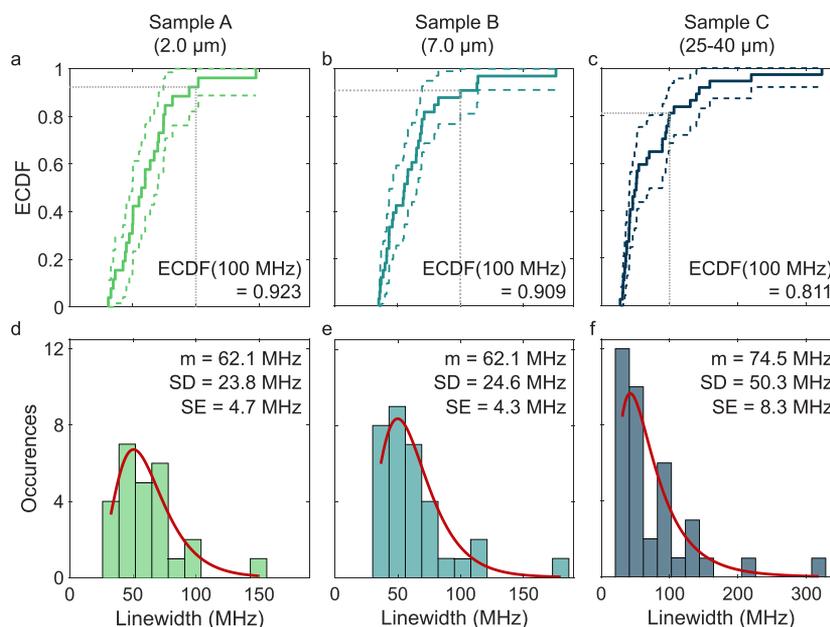


Figure 5. Linewidth statistics of samples A (a,d), B (b,e), and C (c,f), with empirical cumulative distribution functions (ECDFs, a–c) with 95% confidence intervals and linewidth histograms with log-normal fits (d–f). The values of the ECDFs for 100 MHz as well as the means (m), standard deviations (SD), and the standard errors of the mean (SE) are specified.

distributions.²⁶ The corresponding standard errors of the mean (SE) are 4.7, 4.3, and 8.3 MHz, respectively.

A sensible target linewidth is perhaps 100 MHz: in a microcavity, the Purcell effect can increase the homogeneous linewidth to several hundred MHz such that an inhomogeneous broadening of about 100 MHz can be tolerated. The empirical cumulative distribution functions (ECDFs) show that from the measured distributions, one can extract 92.3, 90.9, and 81.1% chances of measuring a linewidth below 100 MHz for samples A, B, and C, respectively. This result emphasizes the excellent optical quality and low-charge-noise environment of the created NV centers. Similar probability values are obtained by using a Bayesian approach for analyzing linewidth distributions;²⁶ with this approach, we determine the probabilities that the next measured line has a linewidth below 100 MHz of 92.3, 92.1, and 78.2% for samples A, B, and C, respectively. With the same approach, we also calculate the probability that the median of the distributions is below this threshold, giving a probability of above 99.9% for each of the three samples. The higher mean linewidth of the NV centers in sample C could be related to the fact that the writing was performed deeper inside the diamond, where it is particularly difficult to avoid spherical aberrations (even with the corrective capability of the objective lens), which means that the vacancies were created over a larger volume compared to the other two samples. Of the 15, 20, and 14 NV centers measured in samples A, B, and C, respectively, 13, 15, and 13 showed clear PLE signals.

CONCLUSION

In conclusion, we have demonstrated that a t-SIL enables laser-induced vacancy creation within bulk diamond with pulse energies as low as 5.8 nJ. There is a large window in pulse energy before the pulse creates irreversible damage. We interpret this wide range of useful pulse energies as a consequence of working in the tunneling breakdown regime for vacancy creation. The vacancies can be created across the full depth of 40 μm bulk diamond samples. Up to an estimated 92.3% of the created NV

centers have an inhomogeneously broadened linewidth below 100 MHz, illustrating that laser writing yields an exceptionally high probability of generating narrow-linewidth NV centers as compared to standard implantation and annealing.^{22,25,26} This metric is crucial for applications based on spin-photon entanglement, as it is greatly beneficial to perform experiments with minimal preselection of NV centers. Importantly, the linewidths presented contain the full inhomogeneous broadening. To the best of our knowledge, this is the lowest charge noise measured to date including the full effects of the off-resonant charge-state repump.^{17,22,31} The narrow linewidths suggest reduced damage in the vicinity of the NV centers compared to other NV creation methods. The t-SIL lowers significantly the threshold pulse energy for vacancy generation. This result points to the feasibility of using a standard Ti:sapphire ultrafast laser without the need for a regenerative amplifier for vacancy creation in diamond.

The main goal of the current work is implementation of the laser-written NV centers into a diamond-cavity system, which in the first stage requires etching of the diamond down to a few micrometers. On account of the strongly reduced lattice damage and improved linewidth statistics compared to other methods, the hope is that after etching the charge noise will remain low. If the low charge noise is maintained, it should be a key step toward generating spin–spin entanglements at higher rates than what has been achieved so far. In a broader context, the high-optical-quality NV centers created through laser writing present a major advancement not only for cavity applications but also in any application requiring a low-charge-noise environment and charge-state-stable color centers.

AUTHOR INFORMATION

Corresponding Author

Viktoria Yurgens – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0001-9550-1286; Email: viktoria.yurgens@unibas.ch

Authors

Josh A. Zuber – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0002-0079-466X

Sigurd Flågan – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0003-0272-7601

Marta De Luca – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0001-6061-3261

Brendan J. Shields – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0003-1937-2763

Ilaria Zardo – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0002-8685-2305

Patrick Maletinsky – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0003-1699-388X

Richard J. Warburton – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; orcid.org/0000-0002-3095-3596

Tomasz Jakubczyk – Department of Physics, University of Basel, CH-4056 Basel, Switzerland; Faculty of Physics, University of Warsaw, 02-093 Warsaw, Poland; orcid.org/0000-0002-8875-4514

Complete contact information is available at:

<https://pubs.acs.org/10.1021/acsp Photonics.1c00274>

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

We acknowledge financial support from the National Centre of Competence in Research (NCCR) Quantum Science and Technology (QSIT), a competence center funded by the Swiss National Science Foundation (SNF) as well as from the EU FET-OPEN Flagship Project ASTERIQS (grant No. 820394), the SNF project grant No. 188521, and the SNF R'Equip grant No. 170741. T.J. acknowledges support from the European Unions Horizon 2020 Research and Innovation Programme under the Marie Skłodowska-Curie grant agreement No. 792853 (Hi-FrED) and support from the Polish National Agency for Academic Exchange under Polish Returns 2019 programme (agreement PPN/PPO/2019/1/00045/U/0001). S.F. acknowledges support from the Initial Training Network (ITN) SpinNANO. We thank Lucas Thiel for skillful coding assistance and Yannik Fontana for reviewing the manuscript.

REFERENCES

- (1) Jelezko, F.; Gaebel, T.; Popa, I.; Domhan, M.; Gruber, A.; Wrachtrup, J. Observation of coherent oscillation of a single nuclear spin and realization of a two-qubit conditional quantum gate. *Phys. Rev. Lett.* **2004**, *93*, 130501.
- (2) Jelezko, F.; Gaebel, T.; Popa, I.; Gruber, A.; Wrachtrup, J. Observation of coherent oscillations in a single electron spin. *Phys. Rev. Lett.* **2004**, *92*, 076401.
- (3) Maze, J. R.; Stanwix, P. L.; Hodges, J. S.; Hong, S.; Taylor, J. M.; Cappellaro, P.; Jiang, L.; Dutt, M. G.; Togan, E.; Zibrov, A.; et al. Nanoscale magnetic sensing with an individual electronic spin in diamond. *Nature* **2008**, *455*, 644–647.
- (4) Rondin, L.; Tetienne, J. P.; Hingant, T.; Roch, J. F.; Maletinsky, P.; Jacques, V. Magnetometry with nitrogen-vacancy defects in diamond. *Rep. Prog. Phys.* **2014**, *77*, 056503.

- (5) Togan, E.; Chu, Y.; Trifonov, A. S.; Jiang, L.; Maze, J.; Childress, L.; Dutt, M. V. G.; Sørensen, A. S.; Hemmer, P. R.; Zibrov, A. S.; Lukin, M. D. Quantum entanglement between an optical photon and a solid-state spin qubit. *Nature* **2010**, *466*, 730–734.

- (6) Hensen, B.; et al. Loophole-free Bell inequality violation using electron spins separated by 1.3 kilometres. *Nature* **2015**, *526*, 682–686.

- (7) Barbour, R. J.; Dalgarno, P. A.; Curran, A.; Nowak, K. M.; Baker, H. J.; Hall, D. R.; Stoltz, N. G.; Petroff, P. M.; Warburton, R. J. A tunable microcavity. *J. Appl. Phys.* **2011**, *110*, 053107.

- (8) Albrecht, R.; Bommer, A.; Deutsch, C.; Reichel, J.; Becher, C. Coupling of a single nitrogen-vacancy center in diamond to a fiber-based microcavity. *Phys. Rev. Lett.* **2013**, *110*, 243602.

- (9) Johnson, S.; Dolan, P. R.; Grange, T.; Trichet, A. A. P.; Hornecker, G.; Chen, Y. C.; Weng, L.; Hughes, G. M.; Watt, A. A. R.; Auffèves, A.; Smith, J. M. Tunable cavity coupling of the zero phonon line of a nitrogen-vacancy defect in diamond. *New J. Phys.* **2015**, *17*, 122003.

- (10) Kaupp, H.; Hümmer, T.; Mader, M.; Schleder, B.; Benedikter, J.; Haussler, P.; Chang, H. C.; Fedder, H.; Hänsch, T. W.; Hunger, D. Purcell-Enhanced Single-Photon Emission from Nitrogen-Vacancy Centers Coupled to a Tunable Microcavity. *Phys. Rev. Appl.* **2016**, *6*, 054010.

- (11) Janitz, E.; Ruf, M.; Dimock, M.; Bourassa, A.; Sankey, J.; Childress, L. Fabry-Perot microcavity for diamond-based photonics. *Phys. Rev. A: At, Mol, Opt. Phys.* **2015**, *92*, 43844.

- (12) Bogdanović, S.; van Dam, S.; Bonato, S.; Coenen, L. C.; Zwerver, A.-M. J.; Hensen, B.; Liddy, M. S. Z.; Fink, T.; Reiserer, A.; Lončar, M.; Hanson, R. Design and low-temperature characterization of a tunable microcavity for diamond-based quantum networks. *Appl. Phys. Lett.* **2017**, *110*, 171103.

- (13) Riedel, D.; Söllner, I.; Shields, B. J.; Starosielec, S.; Appel, P.; Neu, E.; Maletinsky, P.; Warburton, R. J. Deterministic enhancement of coherent photon generation from a nitrogen-vacancy center in ultrapure diamond. *Phys. Rev. X* **2017**, *7*, 031040.

- (14) Janitz, E.; Bhaskar, M. K.; Childress, L. Cavity quantum electrodynamics with color centers in diamond. *Optica* **2020**, *7*, 1232–1252.

- (15) Heupel, J.; Pallmann, M.; Körber, J.; Merz, R.; Kopnarski, M.; Stöhr, R.; Reithmaier, J. P.; Hunger, D.; Popov, C. Fabrication and Characterization of Single-Crystal Diamond Membranes for Quantum Photonics with Tunable Microcavities. *Micromachines* **2020**, *11*, 1080.

- (16) Riedel, D.; Flågan, S.; Maletinsky, P.; Warburton, R. J. Cavity-Enhanced Raman Scattering for In Situ Alignment and Characterization of Solid-State Microcavities. *Phys. Rev. Appl.* **2020**, *13*, 014036.

- (17) Ruf, M.; Ijspeert, M.; Van Dam, S.; De Jong, N.; Van Den Berg, H.; Evers, G.; Hanson, R. Optically coherent nitrogen-vacancy centers in micrometer-thin etched diamond membranes. *Nano Lett.* **2019**, *19*, 3987–3992.

- (18) Maze, J. R.; Gali, A.; Togan, E.; Chu, Y.; Trifonov, A.; Kaxiras, E.; Lukin, M. D. Properties of nitrogen-vacancy centers in diamond: the group theoretic approach. *New J. Phys.* **2011**, *13*, 025025.

- (19) Schmidgall, E. R.; Chakravarthi, S.; Gould, M.; Christen, I. R.; Hestoffer, K.; Hatami, F.; Fu, K.-M. C. Frequency Control of Single Quantum Emitters in Integrated Photonic Circuits. *Nano Lett.* **2018**, *18*, 1175–1179.

- (20) Bourgoïn, J. C.; Massarani, B. Threshold energy for atomic displacement in diamond. *Phys. Rev. B* **1976**, *14*, 3690–3694.

- (21) Campbell, B.; Mainwood, A. Radiation damage of diamond by electron and gamma irradiation. *Phys. Status Solidi A* **2000**, *181*, 99–107.

- (22) Chu, Y.; et al. Coherent optical transitions in implanted nitrogen vacancy centers. *Nano Lett.* **2014**, *14*, 1982–1986.

- (23) Pezzagna, S.; Rogalla, D.; Wildanger, D.; Meijer, J.; Zaitsev, A. Creation and nature of optical centres in diamond for single-photon emission – overview and critical remarks. *New J. Phys.* **2011**, *13*, 035024.

- (24) Appel, P.; Neu, E.; Ganzhorn, M.; Barfuss, A.; Batzer, M.; Gratz, M.; Tschöpe, A.; Maletinsky, P. Fabrication of all diamond scanning probes for nanoscale magnetometry. *Rev. Sci. Instrum.* **2016**, *87*, 063703.

- (25) Van Dam, S. B.; Walsh, M.; Degen, M. J.; Bersin, E.; Mouradian, S. L.; Galiullin, A.; Ruf, M.; Ijspeert, M.; Taminiau, T. H.; Hanson, R.; Englund, D. R. Optical coherence of diamond nitrogen-vacancy centers formed by ion implantation and annealing. *Phys. Rev. B: Condens. Matter Mater. Phys.* **2019**, *99*, 161203.
- (26) Kasperczyk, M.; Zuber, J. A.; Barfuss, A.; Kolbl, J.; Yurgens, V.; Flagan, S.; Jakubczyk, T.; Shields, B.; Warburton, R. J.; Maletinsky, P. Statistically modeling optical linewidths of nitrogen vacancy centers in microstructures. *Phys. Rev. B: Condens. Matter Mater. Phys.* **2020**, *102*, 075312.
- (27) Bernien, C.; Childress, L.; Robledo, L.; Markham, M.; Twitchen, D.; Hanson, R. Two-Photon Quantum Interference from Separate Nitrogen Vacancy Centers in Diamond. *Phys. Rev. Lett.* **2012**, *108*, 043604.
- (28) Sipahigil, A.; Goldman, M. L.; Togan, E.; Chu, Y.; Markham, M.; Twitchen, D. J.; Zibrov, A. S.; Kubanek, A.; Lukin, M. D. Quantum Interference of Single Photons from Remote Nitrogen-Vacancy Centers in Diamond. *Phys. Rev. Lett.* **2012**, *108*, 143601.
- (29) Tamarat, P.; Gaebel, T.; Rabeau, J. R.; Khan, M.; Greentree, A. D.; Wilson, H.; Hollenberg, L. C. L.; Prawer, S.; Hemmer, P.; Jelezko, F.; Wrachtrup, J. Stark shift control of single optical centers in diamond. *Phys. Rev. Lett.* **2006**, *97*, 083002.
- (30) Lekavicius, I.; Oo, T.; Wang, H. Diamond Lamb wave spin-mechanical resonators with optically coherent nitrogen vacancy centers. *J. Appl. Phys.* **2019**, *126*, 214301.
- (31) Chen, Y.-C.; Salter, P. S.; Knauer, S.; Weng, L.; Frangeskou, A. C.; Stephen, C. J.; Ishmael, S. N.; Dolan, P. R.; Johnson, S.; Green, B. L.; Morley, G. W.; Newton, M. E.; Rarity, J. G.; Booth, M. J.; Smith, J. M. Laser writing of coherent colour centres in diamond. *Nat. Photonics* **2017**, *11*, 77–80.
- (32) Buividas, R.; Aharonovich, I.; Seniutinas, G.; Wang, X.; Rapp, L.; Rode, A.; Taniguchi, T.; Juodkakis, S. Photoluminescence from voids created by femtosecond-laser pulses inside cubic-BN. *Opt. Lett.* **2015**, *40*, 5711–5713.
- (33) Hou, S.; Birowosuto, M. D.; Umar, S.; Anicet, M. A.; Tay, R. Y.; Coquet, P.; Tay, B. K.; Wang, H.; Teo, E. H. T. Localized emission from laser-irradiated defects in 2D hexagonal boron nitride. *2D Mater.* **2018**, *5*, 015010.
- (34) Castelletto, S.; Almutairi, A.; Kumagai, K.; Katkus, T.; Hayasaki, Y.; Johnson, B.; Juodkakis, S. Photoluminescence in hexagonal silicon carbide by direct femtosecond laser writing. *Opt. Lett.* **2018**, *43*, 6077–6080.
- (35) Saleem, U.; Birowosuto, D.; Hou, S.; Maurice, A.; Kang, T. B.; Teo, E.; Tchernycheva, M.; Gogneau, N.; Wang, H. Light emission from localised point defects induced in GaN crystal by femtosecond-pulsed laser. *Opt. Mater. Express* **2018**, *8*, 2703–2712.
- (36) Chen, Y.; Salter, P.; Niethammer, M.; Widmann, M.; Kaiser, F.; Nagy, R.; Morioka, N.; Babin, C.; Erlekampf, J.; Berwian, P.; Booth, M.; Wrachtrup, J. Laser writing of scalable single color centers in silicon carbide. *Nano Lett.* **2019**, *19*, 2377–2383.
- (37) Chen, Y.-C.; Griffiths, B.; Weng, L.; Nicley, S.; Ishmael, S. N.; Lekhai, Y. N. D.; Johnson, S.; Stephen, C. J.; Green, B. L.; Morley, G. W.; Newton, M. E.; Booth, M. J.; Salter, P. S.; Smith, J. M. Laser writing of individual atomic defects in a crystal with near-unity yield. *Optica* **2019**, *6*, 662–667.
- (38) Schaffer, C. B.; Brodeur, A.; Mazur, E. Laser-induced breakdown and damage in bulk transparent materials induced by tightly focused femtosecond laser pulses. *Meas. Sci. Technol.* **2001**, *12*, 1784–1794.
- (39) Mao, S. S.; Quéré, F.; Guizard, S.; Mao, X.; Russo, R. E.; Petite, G.; Martin, P. Dynamics of femtosecond laser interactions with dielectrics. *Appl. Phys. A: Mater. Sci. Process.* **2004**, *79*, 1695–1709.
- (40) Gattass, R. R.; Mazur, E. Femtosecond laser micromachining in transparent materials. *Nat. Photonics* **2008**, *2*, 219–225.
- (41) Clark, C. D.; Walker, J.; Ditchburn, R. W. The neutral vacancy in diamond. *Proc. R. Soc. London, Ser. A* **1973**, *334*, 241–257.
- (42) Keldysh, L. V. Ionization in the field of a strong electromagnetic wave. *J. Exp. Theor. Phys.* **1965**, *20*, 1307.
- (43) Konov, V. I. Laser in micro and nanoprocessing of diamond materials. *Laser Photonics Rev.* **2012**, *6*, 739–766.
- (44) Kononenko, V. V.; Vlasov, I. I.; Gololobov, V. M.; Kononenko, T. V.; Semenov, T. A.; Khomich, A. A.; Shershulin, V. A.; Krivobok, V. S.; Konov, V. I. Nitrogen-vacancy defects in diamond produced by femtosecond laser nanoablation technique. *Appl. Phys. Lett.* **2017**, *111*, 081101.
- (45) Serrels, K. A.; Ramsay, E.; Warburton, R. J.; Reid, D. T. Nanoscale optical microscopy in the vectorial focusing regime. *Nat. Photonics* **2008**, *2*, 311–314.
- (46) Serrels, K. A.; Ramsay, E.; Dalgarno, P. A.; Geradot, B.; O'Connor, J. A.; Hadfield, R. H.; Warburton, R. J.; Reid, D. T. Solid immersion lens applications for nanophotonic devices. *J. Nanophotonics* **2008**, *2*, 021854.
- (47) Riedel, D.; Rohner, D.; Ganzhorn, M.; Kaldewey, T.; Appel, P.; Neu, E.; Warburton, R. J.; Maletinsky, P. Low-loss broadband antenna for efficient photon collection from a coherent spin in diamond. *Phys. Rev. Appl.* **2014**, *2*, 064011.
- (48) Tu, R.; Johnson; Cheng, S. *Ray Optics Simulation*. [ricktu288.github.io/ray-optics/](https://github.io/ray-optics/). (Accessed 2020-10-01).
- (49) Kurita, T.; Mineyuki, N.; Shimotsuma, Y.; Fujiwara, M.; Mizuochi, N.; Shimizu, M.; Miura, K. Efficient generation of nitrogen-vacancy center inside diamond with shortening of laser pulse duration. *Appl. Phys. Lett.* **2018**, *113*, 211102.
- (50) Hu, X. J.; Dai, Y. B.; Li, R. B.; Shen, H. S.; He, X. C. The diffusion of vacancies near a diamond (001) surface. *Solid State Commun.* **2002**, *122*, 45–48.
- (51) Onoda, S.; Tatsumi, K.; Haruyama, M.; Teraji, T.; Isoya, J.; Kada, W.; Ohshima, T.; Hanaizumi, O. Diffusion of Vacancies Created by High-Energy Heavy Ion Strike Into Diamond. *Phys. Status Solidi A* **2017**, *214*, 1700160.
- (52) Liu, Y.; Chen, G.; Song, M.; Ci, X.; Wu, B.; Wu, E.; Zeng, H. Fabrication of nitrogen vacancy color centers by femtosecond pulse laser illumination. *Opt. Express* **2013**, *21*, 12843–12848.
- (53) Loudon, R. *The quantum theory of light*, 3rd ed.; Oxford University Press, 2000; pp 72–76.
- (54) Chu, Y.; Markham, M.; Twitchen, D. J.; Lukin, M. D. All-optical control of a single electron spin in diamond. *Phys. Rev. A: At., Mol., Opt. Phys.* **2015**, *91*, No. 021801.