Advances in nano- and microscale NMR spectroscopy using diamond quantum sensors

Robin D. Allert, a Karl D. Briegel a and Dominik B. Bucher a,b

Quantum technologies have seen a rapid developmental surge over the last couple of years. Though often overshadowed by quantum computation, quantum sensors show tremendous potential for widespread applications in chemistry and biology. One system stands out in particular: the nitrogen-vacancy (NV) center in diamond, an atomic-sized sensor allowing the detection of nuclear magnetic resonance (NMR) signals at unprecedented length scales down to a single proton. In this article, we review the fundamentals of NV center-based quantum sensing and its distinct impact on nano- and microscale NMR spectroscopy. Furthermore, we highlight possible future applications of this novel technology ranging from energy research, materials science, to single-cell biology, and discuss the associated challenges of these rapidly developing NMR sensors.

1. Introduction

Since its inception, nuclear magnetic resonance (NMR) spectroscopy has inarguably become one of the workhorses of modern molecular structural analysis. Its significant advantages lie in its non-invasive structure elucidation while being quantitative and usable under standard conditions, making it ideal for synthetic chemistry, molecular biology, and materials science.1 However, the low intrinsic sensitivity of NMR spectroscopy typically restricts its application to macroscopic sample volumes, commonly hundreds of microliters, practically rendering it inapplicable to nano- and microscopic samples. Although novel microfabricated coils have been shown to be a promising avenue towards conventional inductively-detected NMR spectroscopy of sub-nanoliter volumes,2-5 they face challenges to further reduce the sample volume, e.g., difficulties of fabricating the microscopic pick-up coils and the proximity required to detect a signal from an even smaller number of spins.

In the last decade, a novel quantum sensor for magnetic fields has emerged, promising to overcome this sensitivity challenge. This sensor – the nitrogen-vacancy (NV) center – is one of the numerous color defects in diamond.6 It attracted significant interest due to its spin state-dependent photoluminescence and

Robin D. Allert BSc 2020, Technical University of Munich (TU Munich), pursuing his Master’s degree in chemistry focused on physical and organic chemistry supervised by Dominik B. Bucher. His research focuses on facilitating high spectral resolution NV-NMR in microfluidics for chemistry, biology, and biotechnology applications. Furthermore, his interests lie in boosting the sensitivity of these experiments by developing novel and improved NV diamonds for quantum sensing applications.

Karl D. Briegel BSc in chemistry 2018, University of Innsbruck, MSc in chemistry focusing on physical and theoretical chemistry 2020, Technical University of Munich (TU Munich), PhD student 2021 – now, supervised by Dominik B. Bucher. He is working on spatially resolved NV-NMR techniques for applications in chemistry and biotechnology. Apart from this, he is interested in applying methods from quantum optimal control to maximize the sensitivity of NV-NMR experiments.
straightforward spin manipulation enabling its use as a sensor for various physical quantities such as temperature, strain, and particularly interesting for chemists – oscillating magnetic fields, the decisive parameter in NMR spectroscopy.7–9 Furthermore, due to its atomic size, the NV center can be brought into close proximity to the NMR sample, increasing signal strength and, accordingly, enabling NMR spectroscopy down to single cell volumes,10–12 single molecules13–16 and even single spins.17–19

2. The nitrogen-vacancy center

2.1 The structure and formation of NV centers

The nitrogen-vacancy center (see Fig. 1b) is a point defect in diamond consisting of a nitrogen atom substituting a carbon atom (called P1 center) and an adjacent lattice vacancy.6,20 The tetrahedral crystal structure of diamonds allows four distinct NV center orientations. Even though NV centers occur naturally, they are typically produced on purpose in highly purified synthetic diamonds to ensure high homogeneity and controllable properties for their application in quantum sensing.21,22

Though different quantum sensing applications require specialized NV diamonds, the NV center creation follows a general pathway of nitrogen incorporation, vacancy formation, and subsequent high-temperature annealing to form NV centers.23 Generally, two methods of NV center creation for NMR applications prevail: ion implantation and homoepitaxial diamond growth.

Nitrogen atoms are often incorporated into the diamond by ion implantation (see Fig. 1d), where nitrogen ions are accelerated, penetrate the diamond’s surface, and form P1 centers. Here, the ion’s kinetic energy controls the penetration depth and, thus, the later NV center-depth (i.e. the distance from the surface).21,24,25 Furthermore, the total nitrogen ion fluence allows the NV center density fine-tuning. During the implantation process, ions scatter in the lattice, colliding with carbon atoms – subsequently forming lattice vacancies which can diffuse efficiently inside a diamond at temperatures above 800 °C. In addition, the strain induced by the P1 centers causes the vacancies to diffuse towards the nitrogen26 – accordingly forming NV centers upon combination.27 However, the implantation process typically limits the NV center-depth to tens of nanometers due to exceedingly high energies required which cause unsustainable lattice damage.

Additionally, nitrogen-doped epitaxial-overgrowth of seed diamonds by chemical vapor deposition (CVD, see Fig. 1e) allows the production of homogenous NV center-doped diamond layers with a wide range of nitrogen concentrations (several ppb to tens of ppm P1) and thicknesses (ranging from nanometers to hundreds of micrometers).22,28 Likewise, a small fraction of P1 centers (<0.5%) already capture vacancies during the CVD growth due to lattice strain.29 However, the NV center concentration is significantly improved by creating additional vacancies, which can be achieved by high-energy irradiation with, e.g., electrons,30–32 ions,33,34 or lasers.34,35 Like implanted diamonds, subsequent high-temperature annealing will cause the vacancies and P1 centers to form NV centers.

Commonly, NV center-doped CVD grown layers tend to exhibit more promising properties for quantum sensing than implanted samples, such as increased coherence times and increased NV center yields.36,37 Additionally, one significant advantage of CVD growth is the potential to form preferentially aligned NV ensembles along a single crystal axis.38–40 However, CVD growth risks incorporating unwanted defects, e.g., NVH or crystal dislocations.22 Furthermore, nanometer-thick NV center-doped layers by CVD growth are generally more challenging to realize and reproduce due to the complexity of the process.41,42 On the contrary, implanting NV centers is a less complex process allowing for predictable nitrogen densities and depth distributions.41,42 Furthermore, single-ion implanters enable the deterministic placement of single NV centers.43–45

The field of quantum material science has further developed various methods for NV center creation such as high-energy implantation with protons for micron layers,46 nanometer delta-doped layers,47 or the use of high-temperature high-pressure (HPHT) diamonds.37 For an in-depth discussion on color center creation, we would like to refer to an excellent review by Smith et al.48

2.2 Spin and photophysical properties of the NV center

The electronic structure of the NV center is governed by five electrons; where two electrons are provided by the lone pair of the nitrogen atom and three by carbon atoms adjacent to the lattice vacancy forming an electronic system around the vacancy comparable to molecular orbitals with long-lived spin states and well-defined optical transitions.49–51 Several charge variants of the NV center exist; however, only the negatively charged NV− is active for quantum sensing. Thus, our community uses the NV center synonymously with the NV− charge state. The electronic ground state consists of the triplet states $m_s = |0\rangle$ and $|\pm 1\rangle$ with a zero-field splitting of $\sim 2.87 \text{ GHz}$.
the latter are degenerate at zero magnetic field. After applying an external magnetic field $B_0$ along the NV center’s axis, the $m_s = |±1\rangle$ states will split (Zeeman splitting, $2\gamma B_0$ with $\gamma \approx 28$ GHz T$^{-1}$). Crucially, the optical transitions between the ground state and the excited state are spin conserving, meaning that the optical transitions do not change the spin state $m_s$ of the involved electron (see Fig. 1a). Upon excitation of the NV center with light below $\sim 640$ nm (zero phonon line), the NV center has two relaxation pathways: either by fluorescence or a non-radiative pathway through intermediate singlet states which includes intersystem crossing. Notably, the $m_s = |±1\rangle$ excited states have a higher probability of relaxing through the non-radiative channel compared to the $m_s = |0\rangle$ state, leading to two crucial effects: first, the fluorescence rate of the $m_s = |±1\rangle$ states is reduced compared to the $m_s = |0\rangle$ states, and secondly, the optical excitation leads to spin polarization of the NV centers into the $m_s = |0\rangle$ state. These properties provide a convenient route for the spin state’s preparation and read-out. Moreover, the transitions in the optical ground state between the $m_s = |0\rangle$ and the $|±1\rangle$ states can be addressed by microwave (MW) irradiation, permitting coherent manipulation of the NV centers spin state. In summary, the three vital NV center properties are:

1. **Efficient spin state preparation** to start in a well-defined quantum state,
2. **Coherent manipulation of the spin state** for controlled interaction with the environment, and
3. **Optical spin state read-out** to translate the spin state after the interaction with its environment to a measurable parameter.

These properties enable the application of NV centers as fully functional qubits, the basic unit of quantum technology, one of the few operational at room temperature.$^{22}$ While most branches of quantum technology try to isolate their fragile qubits as much as possible from the environment, quantum sensing hinges on the exceptional interaction of the qubit with its environment to perform measurements, e.g., of magnetic fields.

### 3. Quantum sensing

In general, quantum sensing refers to the process of employing a quantum system, often a quantum coherence, to measure a physical quantity – ideally with superior sensitivity, precision, or length scale compared to classical measurements.$^{53}$

#### 3.1 Static magnetic field sensing using NV centers

The fundamental magnetic sensing scheme applies continuous excitation of the NV center with laser light while sweeping the applied microwave frequency. This measurement scheme probes the transition of the $m_s = |0\rangle$ to $m_s = |±1\rangle$ state and, in turn, measures the strength of the applied magnetic field along the NV center’s axis according to the Zeeman splitting ($2\gamma B_0$). At the start of the measurement, the NV center is in the bright $m_s = |0\rangle$ state due to optical polarization. If the MW frequency matches the transition $m_s = |0\rangle \rightarrow |±1\rangle$, the population of the $m_s = |0\rangle$ state decreases, thus the overall fluorescence intensity decreases (see Fig. 1c). Consequently, NV centers can translate magnetic fields into optical signals (optically detected magnetic resonance, ODMR), measured with a photodiode or
camera, with widespread applications.\textsuperscript{8,54,55} However, this scheme can only measure static (dc) or slowly varying (~100 kHz) magnetic fields and, thus, is not applicable to NMR spectroscopy where the detection of alternating (ac) magnetic fields in the MHz range is commonly required.

### 3.2 Alternating magnetic field sensing using NV centers

To sense ac magnetic fields, we utilize well-established pulse sequences from classical NMR; however, rather than applying them to the sample nuclei, we apply the pulse sequence to our NV centers. In other words – NMR signals are detected from the sample nuclei, we apply the pulse sequence to our NV centers to selectively interact with the environment. To enable pulse sequences for NMR detection, a controllable two-level spin system (qubit) is prepared (see Fig. 1a) by applying an external magnetic bias field $B_0$ along the NV center’s axis to separate the $m_s = |\pm 1\rangle$ states. This enables us to selectively address the $m_s = |0\rangle$ state, then a MW pulse sequence is applied to coherently manipulate the spin state of the NV centers – usually to decouple the NV centers from unwanted signals/noise and to enable the NV centers to selectively interact with the environment. To enable pulse sequences for NMR detection, a controllable two-level spin system (qubit) is prepared (see Fig. 1a) by applying an external magnetic bias field $B_0$ along the NV center’s axis to separate the $m_s = |\pm 1\rangle$ states. This enables us to selectively address the $m_s = |0\rangle$ state, then a MW pulse sequence is applied to coherently manipulate the spin state of the NV centers – usually to decouple the NV centers from unwanted signals/noise and to enable the NV centers to selectively interact with the environment.

#### 3.2.1 Hahn echo experiment on the NV center

Most ac sensing pulse sequences are based on the well-known Hahn echo experiment in magnetic resonance spectroscopy. The Hahn echo pulse sequence starts with the NV center in the $m_s = |0\rangle$ state after the initial laser pulse (see Fig. 2c). Then, the first $\pi/2$ pulse ($90^\circ$ pulse) rotates the Bloch vector of the NV center into the equatorial plane of the Bloch sphere (see Fig. 2a). MW pulses control the position of the respective Bloch vector, and the quantum state can be read out optically via its spin-dependent fluorescence; therefore, translating the acquired NMR signal into an optical signal.

Within this article, we would like to introduce briefly the field of quantum sensing pulse sequences and refer to several in-depth reviews on the topic of quantum sensing for further reading.\textsuperscript{53,55-57}

#### 3.2.2 AC sensing with dynamic decoupling sequences

The properties of the NV Hahn echo experiment enable us to measure external ac magnetic fields caused, for example, by the Larmor precession of nuclei within the diamond or from a chemical sample outside of the diamond. The general method can be compared to the well-established electron spin echo envelope modulation (ESEEM) experiments known from classical ESR spectroscopy.\textsuperscript{58,59} Typically, the influence on the Hahn echo will be minor if the free precession time $t$ is much shorter or much longer than the period of the nuclear Larmor frequency (ac magnetic field). However, if the period $T_{\text{NMR}}$ of the ac frequency matches $2t$, the phase accumulation will not cancel. Instead, the phase accumulation of the two free precession periods will add since the sign of the ac magnetic field has changed after the $\pi$ pulse (see Fig. 2d). Thus, upon sweeping the free precession time $t$, a decrease in the fluorescence at the time $t$ corresponding to $T_{\text{NMR}}/2$ can be observed. This forms the basis for the majority of the ac quantum sensing schemes and NV-NMR. The sensitivity and the spectral resolution of this scheme is limited by the coherence time of the NV center.

As already mentioned, the magnetic noise inside of the diamond is mainly caused by other paramagnetic defects, ultimately limiting the coherence time $T_2$ and therefore the sensitivity. Thus, material engineering can advance ac sensitivity by employing high-purity synthetic diamonds (Type IIa) or even isotopically-engineered diamonds. However, another technique for enhancing the sensitivity is the application of multipulse sequences\textsuperscript{60,61} based on the Carr–Purcell–Meiboom–Gill (CPMG) sequences known from conventional NMR spectroscopy. These sequences achieve more effective decoupling of the NV center from the ambient magnetic noise by applying additional $\pi$ pulses on the NV center. Correspondingly, applying these $\pi$ pulse trains will refocus the Bloch vector.
consecutively with every $\pi$ pulse, and effectively prolonging the $T_2$ time, facilitating longer measurement times and, thus, higher sensitivities. Hence, the NV centers coherence time can reach up to milliseconds at room temperature$^{62}$ and hundreds of milliseconds at cryogenic temperatures. After the read-out, the experiment must be restarted, limiting the spectral resolution achievable to a few kHz. However, more sophisticated pulse sequences have been established, extending the practical measurement time, and reaching theoretically arbitrary spectral resolution.$^{10,63–66}$ Interested readers are referred to excellent reviews.$^{53,56,57}$ Further note that reliable information about the quantum state of the NV center can only be obtained by multiple read-outs to gain sufficient statistics (quantum mechanics is intrinsically probabilistic, spin projection noise). Moreover, the read-out fidelity is typically low due to the limited number of detected photons per individual NV center (shot noise).

### 3.3 Detection volume of an NV-NMR experiment

The key advantage of the NV center, compared to macroscopic induction coils, is its atomic size enabling close proximity to the sample nuclei. As a rule of thumb, the NMR signal detected
by an NV center originates from a hemisphere on the diamond’s surface where the radius equals the NV center’s depth from the surface (see Fig. 2b). For nanoscale experiments the NV centers are close (a few nanometers) to the diamond surface, whereas in microscale NV-NMR, micrometer thick NV ensembles are utilized. It is important to note that an NV ensemble effectively acts as a single sensor with an average detection volume equal to the average NV center depth. The detection volume itself determines in which polarization regime (i.e., statistical or Boltzmann nuclear polarization) NMR spectroscopy can be performed – see the Appendix.

4. Nanoscale NMR spectroscopy

4.1 Fundamentals of nanoscale NMR spectroscopy

In a typical nanoscale NV-NMR experiment where the NV center is 2–10 nm below the diamond’s surface, the NV center, therefore, detects a couple of hundreds to thousands of spins (1 nm$^3 = \sim$70 protons in water). On these small length scales, the number of nuclear spins contributing to the NMR signal is not determined by the Boltzmann nuclear polarization (BNP) – but by random spin fluctuations (statistical nuclear polarization). This statistical nuclear polarization scales with one over the square root of the number of nuclear spins and reaches polarizations of up to a few percent resulting in relatively large magnetic signals at nearby NV centers. These signals can be easily detected with dynamic decoupling sequences, however not without a major caveat: nanoscale NV-NMR does not yet reach spectral resolution sufficient for structure elucidation. Molecular diffusion is a substantial issue for liquid nanoscale NMR; water molecules, for example, diffuse through the detection volume of a single NV center in a few nanoseconds, effectively limiting the sensing time and, accordingly, broadens the linewidth (for water a few MHz to GHz). Thus, the diffusion of molecules determines the linewidth achievable in nanoscale NMR, currently restricting it to high viscosity/low diffusion constant samples, such as oils, polymers, or solids. For solids, anisotropic interactions such as dipolar couplings or chemical shift anisotropy induce spectral line broadening. This effect is well known from the field of solid-state NMR spectroscopy typically mitigated by magic angle spinning (MAS). However, MAS is technically challenging to implement for NV-NMR experiments.

4.2 Single NV center experiments

The field of NV-NMR originated from solid-state physics studying single NV centers as nanoscale magnetometers offering a method to measure magnetic fields with sub-nanometer resolution. This field arose only a decade ago and since then has shown remarkable results detecting NMR signals from nanoscopic length scales. We want to highlight a few of them of particular interest for chemistry and biology.

In 2013, two landmark papers from Staudacher et al. and Mamin et al. successfully utilized a single NV center located approximately 7 nm under the diamond’s surface to detect NMR signals from oils and poly(methyl methacrylate) (PMMA). Hereby, the depth of the NV center corresponds to a sensing volume of only five cubic nanometers – comparable to the size of a single mid-sized protein. This work was expanded by Loretz et al., demonstrating the measurement of $^1$H-NMR with an NV center $\sim$2 nm below the surface equivalent to a measurement volume of eight yoctoliter ($8 \times 10^{-24}$ L). Compared to state-of-the-art microcoil NMR ($>0.1$ nL), the required sample volume to generate a detectable signal was therefore reduced by over twelve orders of magnitude.

These ultra-low-volume detections were pushed to the limit by detecting the magnetic resonance of four silicon nuclei in the strong coupling regime and later detecting a single proton. In this regime, the interaction of the detected nuclei with the NV center is greater than the coupling to the surrounding other nuclei constituting an exceptional case for NMR in which the magnetic field of the individual spins is measured rather than the magnetization of the sample. Detection of a single proton has also been achieved by Sushkov et al. by utilizing reporter spins (stable electron spins at the diamond surface) interacting with a proton and subsequently reading out these reporters by double electron–electron resonance (DEER) spectroscopy using a single NV center.

These proof-of-concept experiments were extended into chemically and biologically relevant samples in 2016. Louchchinsky et al. demonstrated the detection of $^3$H-NMR (see Fig. 3c) and $^{13}$C-NMR (see Fig. 3d) signals form a single protein (see Fig. 3a). The group utilized EDC/NHS crosslinking to covalently bind ubiquitin to the diamond surface (see Fig. 3b) and achieved co-localization of single proteins and single NV centers. Although highly impressive from a physics perspective (see Fig. 3d), the practical applications are limited due to broad resonance lines in the dried solid-state they used for their measurement.

Nevertheless, utilizing NMR to its full potential requires high spectral resolution – two essential parameters need to be resolved for structure elucidation: the $J$-coupling and the chemical shift. Neighboring nuclear spins cause $J$-coupling (typically in the range of 0.1–12 Hz for protons in small organic molecules independent of the magnetic bias field $B_0$), mediated through chemical bonds, that offers information of the type, orientation, and number of close neighboring nuclei. Chemical shift is induced by the electron density of chemical groups close to the nuclei causing miniscule (ppm) magnetic field differences at the nuclear spin’s location, modifying the local magnetic field and thus causing a relative shift of the resonance frequency with respect to the bias field $B_0$. Therefore, increasing the magnetic field $B_0$ is beneficial for chemical shift resolution. Consequently, NMR spectroscopy is preferably performed at high magnetic fields $B_0$; the implications for these high magnetic fields in NV-NMR experiments will be discussed in Section 5.4. Aslam et al. successfully demonstrated chemical shift resolution at the nanoscale utilizing a 3 Tesla superconducting magnet by using an advanced quantum sensing protocol which extended the practical relaxation time of the NV sensor by several orders of magnitude. This pulse...
sequence utilizes the intrinsic nitrogen nuclear spin of the NV center as a quantum memory, thus improving the theoretically achievable linewidth resolution to 0.1 mHz or 0.01 ppb at 3 T for nanoscale NMR, in line with a modern conventional NMR spectrometer. However, as discussed earlier, the nanoscopic sample itself poses a fundamental limitation on the linewidth. Aslam et al. used highly viscous fluids as samples to minimize the effect of diffusion on the linewidth, which enabled them to measure chemical shift resolved $^1$H and $^{31}$P-spectra with a linewidth of 1.3 ppm and 1.4 ppm.

One promising route towards improving chemical information despite broad resonances in nanoscale NV-NMR is the detection of nuclear quadrupolar spins, i.e., spins with a spin quantum number greater than $S = 1/2$. The quadrupolar splitting is mediated by the interaction of the quadrupolar moment of the nucleus with the electric field gradient of the local electronic structure. These splittings are typically large (tens of kHz to several tens of MHz) and highly sensitive to the molecular structure. Lovchinsky et al.$^{79}$ detected quadrupolar resonances of $^{11}$B and $^{14}$N of atomically thin hexagonal boron nitride (hBN). The authors observed a dependence of the quadrupolar resonance with different layer thicknesses, paving the way to probe local structures in two-dimensional materials.

4.3 Shallow NV ensembles experiments

While single NV centers have superior spatial resolution, there are certain drawbacks including tedious co-localization of the sample and a suitable NV center, intrinsically inhomogeneous properties, and instability of single, near-surface NV centers.$^{81,82}$ Furthermore, the detection of single photons make these experiments susceptible to background fluorescence. Consequently, the overall experimental complexity of single NV center experiments with long averaging times and background fluorescence results in a low throughput for troubleshooting and/or detection of large parameter spaces. In our opinion, these drawbacks render single NV center experiments so far difficult for applications in chemistry or biology. A straightforward method of effectively increasing the system’s robustness, usability, and sensitivity is to employ an ensemble of NV centers close to the diamond’s surface with the trade-off of losing the nanometer-scale spatial resolution. Instead, the diameter of the laser excitation – typically a few microns – and the depth of the NV centers define the measurement volume. This approach lends itself to simpler, robust, and highly sensitive setups, allowing easy NV-NMR integration into chemical or biological research.$^{84}$

DeVience et al.$^{75}$ demonstrated simultaneous NV-NMR of multiple nuclei ($^1$H, $^{19}$F, and $^{31}$P) with shallow NV ensembles and single NV centers. The ensemble approach circumvents the tedious search for a suitable NV center and allows for widefield magnetic resonance imaging (MRI) at the nanoscale imaged by a charge-coupled device (CCD) camera with a lateral resolution of ~500 nm and a field of view of 50 μm. Nanoscale imaging, where each pixel corresponds to an NMR spectrum, is one of the crucial benefits of the optical read-out of NV centers. The widefield NMR imaging approach was further expanded by Ziem et al.$^{85}$ by implementing optimal control sequences for the NV sensors, allowing for more robust spin control even with inhomogeneous MW driving fields over the field of view. They applied their optimized pulses in magnetic resonance imaging of fluorine-containing thin films of 1.2 nm thickness resulting in an effective signal from ~120 nuclei per single NV center. Optimal control sequences are already applied in conventional NMR spectroscopy$^{86-88}$ as well as in NV-based sensing$^{89-91}$ demonstrating great potential to further increase the sensitivity and robustness of the sensing schemes for NV-NMR spectroscopy.

So far, previous work detected the NMR signals directly on top of the diamond from either bulk samples, oils, or samples directly tethered to the diamond’s surface. However, we envision the strength of NV-NMR in probing interfaces and surfaces...
beyond the diamond’s intrinsic surface with applications especially for chemistry, which is one focus of our group.

Interfacial and surface properties are vital for numerous material functions and applications in catalysis, energy conversion, or bioanalytics, especially under chemical-relevant conditions. However, state-of-the-art surface analysis techniques require ultra-high vacuum, intense synchrotron radiation, or femtosecond lasers. Even then, chemical reaction kinetics and molecular dynamics are ultimately challenging to measure under ambient conditions. Liu et al. recently demonstrated the application of NV-NMR in combination with atomic layer deposition (ALD) to study self-assembled monolayers (SAM) under ambient conditions (see Fig. 4a). ALD is a technique that enables the surface deposition of thin films ranging from pure metals, common polymers (utilizing molecular layer deposition, MLD), or metal oxides. In Liu’s study, a one-nanometer thick amorphous aluminum oxide layer was deposited, allowing them to probe the chemical modification of this layer with phosphonate chemistry during the formation of various self-assembled monolayers from organic compounds containing fluorine and phosphorus. The surface NMR technique detected both nuclei from the SAM-modified Al₂O₃ layer, providing chemical information with femtomole sensitivity with few tens of micron spatial resolution (see Fig. 4b), allowing for quantitative studies over the millimeter-sized diamond chip. Additional information on the SAM was obtained by comparing the linewidth of the fluorine signal (see Fig. 4c). As discussed previously, in solid-state NMR spectroscopy the linewidth is predominantly determined by dipolar coupling, which can be minimized by sample motion. Thus, fluorine located at the end of the SAM chain exhibits smaller linewidths due to their greater rotational freedom. Likewise, the line broadening effect can be used to study the binding effect of the phosphonate group to the Al₂O₃ support, where the Al₂O₃-bound moiety had significantly broader linewidths than drop-casted samples without a support layer, indicating chemical binding. In distinction to other analytical methods for surfaces, surface NV-NMR allows time-resolved studies under chemical-relevant conditions enabling observation of the real-time phosphonate-binding to the Al₂O₃ support at the solid-liquid interface (see Fig. 4d). Therefore, NV-NMR enables direct detection of the reaction kinetics at the solid-liquid interface by measuring the NMR signal as a function of time, demonstrating faster SAM formation utilizing higher substrate concentrations.

4.4 Outlook

Even though single NV center studies offer superior spatial resolution and have reached the absolute detection limit of

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Fig. 4 Surface NMR using shallow NV ensembles for probing chemistry at the interfaces. (a) Schematic of the surface NV-NMR experiments on a functionalized aluminum oxide (Al₂O₃) layer utilizing shallow NV ensembles, which measure the statistical polarization of fluorine and phosphorus nuclei from a self-assembled monolayer (SAM) of phosphonates. (b) NV-NMR allows for spatial probing (imaging) of the SAM layer’s homogeneity at several different measurement spots. (c) Chemical information contained in the signal linewidth originating from the SAM can be detected, indicating local molecular dynamics and potentially the binding of the molecules to the aluminum oxide layer. (d) NV-NMR allows for time-resolved in situ studies of the monolayer formation under chemical relevant conditions observing the phosphonate-binding to the support structure, therefore, demonstrating different formation kinetics at various phosphonate concentrations. All figures adapted from K. S. Liu et al., Proc. Natl. Acad. Sci. U. S. A., 2022, 119, e2111607119 with permission from “Copyright (2022) National Academy of Science”.

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single spins, some drawbacks must be overcome to establish them as a general method such as sample and NV center co-localization. Over the past couple of years scanning NV experiments for magnetic field sensing have become a standard tool in solid-state physics and are commercially available. These scanning-probe experiments contain a scanning tip etched from high-purity diamond and include a single NV center. While typically used in nanoscale magnetic imaging, these microscopes could be adapted for NMR imaging—potentially solving the co-localization by bringing single NV centers to the sample allowing for nanoscale magnetic resonance imaging. In a similar fashion, the sample itself can be mounted on an AFM-like tip circumventing the colocalization issue. These tip-based methods, however, face the potential challenge of increased distance between the sample and the respective NV center, reducing the signal strength. Another promising approach, NV-NMR with functionalized nanodiamonds containing NV centers has been successfully demonstrated bringing the NMR detector into the sample itself and, thus, opening the route for NMR inside cells.

By the same token, shallow NV ensembles for NMR spectroscopy have demonstrated their potential in surface sciences, due to the simple setup and robust measurements. Here, we envision a bright future for NV-NMR using NV centers as a novel analytical tool for material, catalysts, and energy research, ideally under in operando conditions.

A recent study has expanded the aluminum oxide platform for surface NMR demonstrated by Liu et al. including additional binding moieties such as biotin or azides by multiple follow-up functionalization of the Al₂O₃ support. This approach enables the efficient coupling of biomolecules such as proteins or DNA to the aluminum oxide deposited on the diamond’s surface and expands the prospect of surface NV-NMR for biomolecules or biochemistry.

For all nanoscale NV-NMR experiments molecular structural information is presently limited by its diffusion-limited linewidth in liquid samples and dipolar coupling in solid-state samples to several kHz. Magic angle spinning known from classical NMR might solve the latter with significant engineering challenges. Furthermore, confinement of liquid samples for example in nanofluidics may restrict diffusion potentially enabling high spectral resolution. Yet, recent results indicate that the diffusion can lead to sharp-peak spectral lines which could enable improved resolution. Another promising way to gain molecular information is the detection of quadrupolar nuclei (such as ²H, ¹¹B, ¹⁴N, …), as discussed before.

5. Microscale NMR spectroscopy

5.1 Fundamentals of microscale NMR spectroscopy

While nanoscale NV-NMR greatly benefits from statistical polarization up to several percent, its diffusion-limited linewidth impedes its application in structural elucidation so far—one of the strong suits of NMR spectroscopy. Microscale NMR using dense NV center-doped diamond layers with several micron thicknesses overcomes this issue by detecting the Boltzmann polarization. Effectively, microscale NV-NMR trades signal strength for spectral resolution. In this case, each equivalent nuclear spin of the sample behaves identically, and molecular diffusion does not limit the linewidth anymore. These experiments resemble conventional NMR experiments where a free nuclear precession (FNP, see the Appendix) is induced by applying a π/2 pulse on the sample nuclei resulting in a coherent NMR signal.

5.2 High resolution microscale NV-NMR spectroscopy

These signals are typically detected by the so-called coherently averaged synchronized read-out (CASR) sequence, meaning that dynamically decoupled sequences are synchronized to the detection signal (e.g., the FNP, see Appendix), allowing for arbitrary long measurement times and thus high spectral resolution. Recent works have demonstrated Hz-linewidths resolving J-couplings and chemical shifts from chemical samples in picoliter volumes using CASR or adapted sequences.

Ultimately, the magnetic field uniformity and stability limited the spectral resolution of these experiments. In addition to the differences in linewidth and polarization, the sample geometry has a significant impact on the signal strength in microscale NV-NMR in contrast to the nanoscale. Similarly, the NMR signal strongly depends on the NV centers orientation with respect to the diamond’s surface—establishing a path for further improvements for the microscale NV-NMR setups. These effects have been investigated in great detail by Bruckmaier et al.

A first principle study by Glenn et al. has demonstrated ¹H-NMR spectra resolving, for example, the distinct ¹H–³¹P J-coupling from trimethyl phosphate (see Fig. 5a and b) and the chemical shift of protons in xylene (see Fig. 5c) from a ten picoliter measurement volume—the volume of a single mammalian cell. Furthermore, even though long measurement times (~1 h) were required to achieve these spectra by solely measuring Boltzmann polarization, this study accomplished similar detection limits (LOD) with a significantly lower sample volume than microscale inductive NMR (see Fig. 5d). However, measuring Boltzmann polarization with NV-NMR is a double-edged sword: while the polarization and, thus, the detectable NMR signal and the chemical information benefits strongly from increased magnetic fields, NV center-based quantum sensing becomes more and more challenging. The dynamic decoupling sequences work most efficiently in the low MHz regime; fundamentally limited by the minimal achievable π pulse duration. Therefore, higher magnetic fields, i.e., higher NV resonance frequencies result in experimental complexity and raises the cost of the microwave equipment for the NV-spin control. For that reason, current microscale experiments have so far only been performed below 0.1 T. The impact of high magnetic fields on NV-NMR will be discussed in Section 5.4.

5.3 Hyperpolarization enhanced microscale NV-NMR experiments

To increase the concentration sensitivity, hyperpolarization methods known from conventional NMR or MRI have been...
successfully applied. First, in a brute-force hyperpolarization approach, the nuclear spin sample is pre-polarized at high magnetic fields, shuttled to the spectrometer then measured at low magnetic fields. Smits et al.\textsuperscript{105} used this method to detect $^1$H-NMR and $^1$H–$^{19}$F-heteronuclear correlated spectroscopy (COSY) of 1,4-difluorobenzene by pre-polarizing the sample at 1.5 T and subsequently measuring it at 13 mT, achieving one ppm linewidths.

Furthermore, microscale NV-NMR can be combined with auxiliary \textit{in situ} hyperpolarization techniques known from NMR such as parahydrogen-induced hyperpolarization (PHIP) or Overhauser dynamic nuclear polarization (DNP).\textsuperscript{107} Overhauser DNP specifically stands out for its ease of implementation in NV-NMR, providing signal enhancements of up to two orders of magnitude for $^1$H-NMR and up to three orders of magnitude for $^{13}$C nuclei.\textsuperscript{108} This method requires the addition of stable radicals into the solution, typically a few mmol L$^{-1}$ of TEMPO or its derivatives, which allows the transfer of Boltzmann polarization of the unpaired electron spin ($\approx$ three orders of magnitude higher than $^1$H polarization) to the nuclear spins of the sample by irradiating the sample with microwaves at the electron transition frequency. While integrating the Overhauser DNP in conventional NMR is usually not trivial because of the required microwave frequencies electronics for driving the electron transition,\textsuperscript{109} this is straightforward in NV-NMR because it already incorporates the necessary Gigahertz electronics for the NV spin control. Furthermore, Overhauser DNP has been shown to work best at low magnetic fields,\textsuperscript{110} facilitating NV-NMR. For example, Bucher \textit{et al.}\textsuperscript{11} showed femtomolar sensitivity ($\approx$ 5 mM concentrations) in a ~ ten picoliter volume.
Correspondingly, they demonstrated enhancement factors between 80–230 times for a broad range of small molecules (Fig. 5e). Nevertheless, while straightforward to implement, Overhauser DNP has some adjoined disadvantages: the enhancement factor varies significantly between molecules and even nuclei within the same molecule, complicating NMR studies. Moreover, the paramagnetic radicals cause line broadening, which may hamper the desired spectral resolution, and the Boltzmann polarization of the electron limits the maximum achievable enhancement.

Higher enhancements have been achieved via PHIP, where the nuclear singlet state of parahydrogen is used as a polarization source. PHIP allows hyperpolarized molecules to be prepared by either hydrogenation reactions of unsaturated precursors (hydrogenative PHIP using PASADENA113 or ALTADENA114 conditions) or by non-hydrogenative via an exchange process where the substrate and the parahydrogen exchange on the same metal center via signal amplification by reversible exchange (SABRE). Arunkumar et al.12 combined the latter technique with microscale NV-NMR to achieve 0.5% proton spin polarization at 6.6 T, improving five orders of magnitude over Boltzmann polarization. Accordingly, this method facilitated the measurement of small-molecule samples (e.g., N-labeled pyridine and nicotinamide) with concentrations as low as one millimolar from a ten picoliter volume, corresponding to 10 fmol, after 300 seconds of averaging time. While conventional NMR and MRI have shown a wide range of molecules polarizable with PHIP-based techniques, allowing for in vivo pH sensing or monitoring metabolites, PHIP is not universally applicable in chemistry or biology. However, the combination with the small volume detection of NV-NMR can open the field to single-cell analysis.

5.4 Outlook

Higher Boltzmann polarization is one of the primary motivations to reach for higher magnetic fields but not the only one; spectral resolution also dramatically benefits from higher magnetic fields splitting the NMR signals and, thus, allowing for better spectral resolution. So far, all NV-NMR experiments at the microscale have been performed below 0.1 T. However, magnetic fields below 0.35 T are an unfortunate regime for chemical analysis where the chemical shift and the J-coupling are at a similar magnitude (several Hz for protons). This “overlapping” of J-coupling and chemical shift induces strong couplings and second-order effects, which in turn cause the resonance lines to split at higher multiplicities than observed in typical high-field NMR. These effects can obscure the usually easy to interpret NMR spectra.

However, the NV center’s resonance frequency at which the pulse sequence is performed and the respective NV center’s π pulses scale with the magnetic bias field $B_0$. While the NV center’s resonance frequency scales with $\sim 28$ GHz $T^{-1}$, the π pulses must be short enough to facilitate the high sample Larmor frequencies (tens of MHz) with dynamic decoupling sequences, i.e., significantly shorter than half the NMR frequency period. These form exceedingly difficult technical challenges requiring higher microwave frequencies and powers, which might cause among other things dielectric sample heating and demand more expensive equipment. However, ESR spectroscopy has solved some of these technical hurdles by establishing microwave resonators and amplifiers in the X band (8–12 GHz) and the K$_\alpha$ band (26–40 GHz) corresponding to an NV-NMR range of approximately 0.2 T to 1.5 T. Likewise, magnets with sufficient homogeneity and stability in the sub-ppm range are widely available at these magnetic fields. Reaching these magnetic fields is one of the major goals of our community and would enable NMR spectra equivalent to modern commercial benchtop NMR devices, however, with significantly lower sample volumes.

NV-NMR is well suited for microscale measurements. Thus, further integration with microfluidics will provide an optimal match of the sample and measurement volumes. Chip-based microfluidics or Lab-on-a-Chip is an established platform for chemical and biological applications. Its benefits hinge primarily on system miniaturization, such as well-understood mass and heat transport, as well as parallel processing, monitoring assays, and small amounts of reactants needed. Current developments aim to reduce sample volumes further down to the picoliter regime. While microfluidics can mimic most other established chemical or biological assay operations, e.g., mixing, incubation, or liquid–liquid extractions, its analytics is limited to fluorescence spectroscopy, absorption spectroscopy, or mass spectrometry. However, the spectroscopic methods do not offer structural information and require either fluorescent samples or often complicated coupled reactions in the case of fluorescence spectroscopy. While mass spectrometry does offer structural information, it destroys the sample and is not generally applicable for every sample and solvent type. NV-NMR could overcome these limitations by offering non-invasive measurements of molecular structural information. Furthermore, NV-NMR could be used to measure parameters such as molecular diffusion, flow rate, pH, or temperature inside the microfluidics, establishing a new analytical platform for Lab-on-a-chip applications.

Likewise, improvements in sensitivity and spectral resolution could enable NMR spectroscopy of small molecules and proteins in single cells with potential applications ranging from single-cell metabolomics to NMR fingerprinting of tumors. Furthermore, the optical read-out of NV-NMR allows for spatial resolution, which can be visualized as “scanning microscopy” by moving the laser excitation and thus the measurement volume. Combined with correlative optical microscopy, this would enable MRI measurements of tissues or biofilms.

6. Hyperpolarization using NV centers

So far, we have only discussed the NV center as a novel NMR sensor. However, the NV center itself has been proposed as a polarization source; optical pumping can rapidly (~μs) and efficiently (>80%) polarize the NV center’s electron spin. The resulting non-equilibrium polarization could be transferred to...
the chemical sample similar to classical DNP. However, unlike established DNP, there is no need for added paramagnetic radicals or cryogenic temperatures. Especially NV center-doped nanodiamonds as a polarization source for classical NMR demonstrate promising results and have been realized using small form factor setups at room temperature, which could facilitate the

<table>
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<th>Regime</th>
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<tr>
<td>Nanoscale NV-NMR</td>
<td>Surfaces and interfaces</td>
<td>Battery research, electrochemistry catalysis, surface chemistry, bioanalytics, thin films, 2D materials, ...</td>
<td>Limited spectral resolution</td>
<td>Nuclear quadrupolar resonance (NQR) spectroscopy, Novel pulse sequences enabling detection at higher magnetic fields, Nanofluidics sample confinement for diffusion restriction, Magic angle spinning, High magnetic fields (e.g., 3 T) in combination with advanced quantum sensing schemes</td>
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<td>Single molecules</td>
<td>Structural biology (i.e., protein or DNA folding), catalysis, trace analysis, ...</td>
<td>Sample co-localization, NV charge state instability, leading to photoionization of the NV-centers, Limited throughput, Limited NV center homogeneity</td>
<td>Co-doping with charge stabilizing electron donors, NV-NMR, Nanofluidics, surface modifications, scanning NV-NMR</td>
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<td>Microscale NV-NMR</td>
<td>Single cells</td>
<td>Single cell biology, metabolomics, tumor detection, directed evolution, ...</td>
<td>Limited concentration sensitivity, Limited spectral resolution, In vivo conditions (medium circulation, cell movement, ...)</td>
<td>Apply hyperpolarization methods such as Overhauser DNP, PHP, dDNP, CIDNP, Improved NV read-out for sensitivity enhancement, Increased magnetic fields (&gt;0.35 T), Ultralow-field NMR in combination with hyperpolarization, Novel pulse sequences enabling detection at higher magnetic fields, Microfluidic devices for cell cultures such as cell traps or nano titer arrays</td>
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<tr>
<td>Lab-on-a-chip</td>
<td>Nanomol chemistry, microdroplet screening, organ-on-a-chip, point-of-care testing, drug screening, ...</td>
<td>Sample heating (microwave and laser heating), Single cell statistics</td>
<td>Limited concentration sensitivity, Limited spectral resolution, In vivo conditions (medium circulation, cell movement, ...)</td>
<td>Apply hyperpolarization methods such as Overhauser DNP, PHP, dDNP, CIDNP, Improved NV read-out for sensitivity enhancement, Increase magnetic fields (&gt;0.35 T), Ultralow-field NMR in combination with hyperpolarization, Novel pulse sequences enabling detection at higher magnetic fields, Microfluidic devices for cell cultures such as cell traps or nano titer arrays</td>
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<tr>
<td>NV-Hyperpolarization</td>
<td>Easy, low-cost, and efficient hyperpolarization for chemical analytics and diagnostics, ...</td>
<td>Number of NV spins to sample spin ratio</td>
<td>Material engineering to increase the NV spin density, Nanostructuring of diamond to enhance the surface area, Nanodiamonds containing NV centers</td>
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practical translation of this technology. Therefore, this method of hyperpolarization could enable easy, low-cost, and efficient hyperpolarization for chemical analytics.132

Furthermore, the combination of NV center-based hyperpolarization with NV-detected NMR could enable applications requiring in situ hyperpolarization like single-cell NMR or NMR of nuclei with low gyromagnetic ratios. So far, only initial proof-of-principle studies have been conducted, demonstrating that polarization from single NV centers139–141 and shallow NV ensembles can be transferred to spins outside of the diamond.142,143 Nevertheless, there has not been a conclusive study demonstrating the subsequent detection of elevated polarization levels of sample spins outside of the diamond. Theoretical evaluations have proposed modest enhancements for microscale NV-NMR by one or two orders of magnitude with significant improvements by micro-structuring the diamond surface. In contrast, nanoscale NV-NMR is unlikely to benefit from this technology at the current stage due to the considerable statistical polarization.144

7. Conclusions

The NV center in diamond has demonstrated its capabilities as a detector for nano- and microscale NMR spectroscopy. Researchers have realized impressive nanoscale NMR measurements of two-dimensional materials,79 single proteins,13 and even single protons.18,19 However, low spectral resolution of nanoscale NV-NMR has limited its applications so far. Since the technique is still in its infancy, future developments such as nuclear quadrupolar spectroscopy79 or liquid sample confinement72 will very likely overcome this drawback (see Table 1). Nevertheless, nanoscale NV-NMR employing shallow NV ensembles has shown its ability to measure NMR signals originating from surfaces and interfaces under chemical-relevant conditions expanding the toolbox of modern chemical analytics with widespread applications in catalysis, electrochemistry, and material research.77

Microscale NV-NMR has made notable progress towards its application in single-cell studies or chemical analytics in microfluidics, demonstrating high spectral resolution and high spin sensitivity.10,66 Furthermore, it can be readily interfaced with established NMR techniques such as hyperpolarization,11,12,105 2D-NMR pulse sequences,105 or different nuclei,105 enabling structural and quantitative information at length scales of a single-cell. Additionally, microscale NV-NMR will benefit significantly from increased magnetic fields in the upcoming years, allowing for chemical shift resolution and sensitivity equivalent to commercial benchtop NMR systems – but with picoliter volumes. We expect considerable progress in the field soon, establishing NV-NMR as a standard tool for chemistry and life sciences.

Author contributions


Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this feature article.

Appendix

Origin of the NMR signal in nano- and microscale NV-NMR experiments

Microscale NV-NMR spectroscopy typically detects Boltzmann nuclear polarization – well-known from conventional NMR – where only minuscule fractions of the sample spins are polarized in the external magnetic field $B_0$ and contribute to the overall sample magnetization $M_S$ (see Fig. 6b). This fraction is determined by the Boltzmann distribution and amounts, e.g., to only ~100 parts per million for a 28 Tesla field and at room temperature. Importantly, nanoscale NV-NMR experiments detect only a small number of spins (e.g., 10 000) – here statistical fluctuations of spin orientations lead to an incomplete cancellation of the magnetic moments; thus resulting in an effective spin polarization of the chemical sample – called statistical nuclear polarization (SNP, see Fig. 6a).68,69 Müller et al.70 provide an intuitive picture of SNP based on a stochastic signal arising from a repeated quantized measurement of a nuclear spin ensemble: if a single spin ($\mathcal{S} = 1/2$) is measured, its transverse angular momentum will result in $-h/2$ for half the time and $+h/2$ respectively for the other half. For two spins, the possible outcomes would be $\pm h$ and 0. Therefore, for vast

| Table 2 Comparison of statistical and Boltzmann nuclear polarization |
|---------------------|---------------------|
| **Nanoscale NMR**   | **Microscale NMR**  |
| Number of spins $N$ | SNP $\propto \sqrt{1/N}$ | Independent |
| Magnetic field $B_0$ | Independent | BNP $\propto B_0$ |
| Temperature $T$     | Independent | BNP $\propto 1/T$ |
| Achievable polarization | Up to several % | $e.g., \sim 100$ ppm at 28 $T$ and 300 K |
| Measured signal     | No sample spin excitation necessary | Active sample spin excitation required ($\pi/2$ pulse) |
|                     | due to the detection of incoherent noise signals; repetition time not limited by sample $T_1$ | to induce coherent free nuclear precession (FNP); repetition time limited by sample $T_1$ |
numbers of spins ($N$), the probability would approximate a Gaussian distribution with a standard deviation $\sigma_{M_z}$ of $+\hbar\sqrt{N}/2$. While the average of all measurements would result in 0, i.e., cancelation of the signal, the standard deviation $\sigma_{M_z}$ of the measured values is a nonzero result. Consequently, nanoscale NMR detects the variance ($\sigma_{M_z}^2$) of the magnetization rather than the sample magnetization $M_z$ (see the inset of Fig. 6c).

The statistical polarization scales with the square root of the inverse number of spins and can reach several percent polarization for a small number of spins (see Fig. 6d), while Boltzmann polarization benefits from higher magnetic fields and lower temperatures (see Table 2). Importantly, SNP detection does not require active excitation of the nuclear spins nor waiting to reach equilibrium ($T_1$ time) due to the detection of incoherent noise signals. However, it comes with some drawbacks, such as diffusion limited linewidths, discussed in the main text. Furthermore, we want to highlight the publication by Schwartz et al. examining the differences and applications of SNP and BNP in great detail.71

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