Background-free dual-mode optical and $^{13}$C magnetic resonance imaging in diamond particles

Xudong Lv$^{a, b, c}$, Jeffrey H. Walton$^{d, e}$, Emanuel Druga$^{a, f}$, Fei Wang$^{g}$, Alessandra Aguilar$^{a, h}$, Tommy McKnelly$^{a, i}$, Raffi Nazaryan$^{a, j}$, Fanglin Linda Liu$^{a, k}$, Lan Wu$^{a, l}$, Olga Shenderova$^{a, m}$, Daniel B. Vigneron$^{a, n}$, Carlos A. Meriles$^{a, o}$, Jeffrey A. Reimer$^{b, p}$, Alexander Pines$^{a, q, r}$, and Ashok Ajoy$^{a, s}$

$^{a}$Department of Chemistry, University of California, Berkeley, CA 94720; $^{b}$Nuclear Magnetic Resonance Facility, University of California, Davis, CA 95616; $^{c}$Department of Electrical Engineering and Computer Sciences, University of California, Berkeley, CA 94720; $^{d}$Adamas Nanotechnologies, Inc., Raleigh, NC 27617; $^{e}$Department of Radiology and Biomedical Imaging, University of California, San Francisco, CA 94158; $^{f}$Department of Physics, City University of New York–City College of New York, New York, NY 10031; $^{g}$City University of New York–City College of New York, New York, NY 10031; $^{h}$Department of Chemical and Biomolecular Engineering, Lawrence Berkeley National Laboratory, University of California, Berkeley, CA 94720; and $^{i}$Materials Science Division, Lawrence Berkeley National Laboratory, University of California, Berkeley, CA 94720

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Multimodal imaging—the ability to acquire images of an object through more than one imaging mode simultaneously—has opened additional perspectives in areas ranging from astronomy to medicine. In this paper, we report progress toward combining optical and magnetic resonance (MR) imaging in such a “dual” imaging mode. They are attractive in combination because they offer complementary advantages of resolution and speed, especially in the context of imaging in scattering environments. Our approach relies on a specific material platform, microdiamond particles hosting nitrogen vacancy (NV) defect centers that fluoresces brightly under optical excitation and simultaneously “hyperpolarize” lattice $^{13}$C nuclei, making them bright under MR imaging. We highlight advantages of dual-mode optical and MR imaging in allowing background-free particle imaging and describe regimes in which either mode can enhance the other. Leveraging the fact that the two imaging modes proceed in Fourier-reciprocal domains (real and k-space), we propose a sampling protocol that accelerates image reconstruction in sparse-imaging scenarios. Our work suggests interesting possibilities for the simultaneous optical and low-field MR imaging of targeted diamond nanoparticles.

In the quest toward high signal-to-noise (SN) imaging, significant power can be brought to bear by multimodal or multimessenger techniques (1, 2). They entail capturing an object through more than one imaging mode simultaneously, often at widely disparate wavelengths. Exploiting correlations between the different modes portends approaches such as Kalman filtering (3) that can deliver noise or background suppression. Furthermore, under appropriate conditions these correlations can engender additional image sampling and reconstruction strategies to accelerate image acquisition.

In this paper, we consider whether such methods could be applied to combine together MRI and optical imaging (4, 5). These two modes offer diametrically complementary advantages. Visible-wavelength optics are fast and cheap and can image at high resolution, yet often suffer from scattering, attenuation, and aberration distortions while imaging through real media. MRI, on the other hand, is noninvasive and scattering-free, is fully three dimensional, and can be chemically functional; yet it is slow, suffers from weak signals, and offers poor spatial resolution (millimeter level). Notably, optical and magnetic resonance (MR) imaging are carried out in Fourier-reciprocal spaces (x- and k-space). This redundancy makes a combined modality attractive—not only are there complementary advantages in sensitivity and resolution to be gained, but also it opens possibilities to access hybrid acquisition strategies that sample both real space and k-space simultaneously to yield image acceleration.

Here we illustrate proof-of-concept demonstrations of dual-mode optical and $^{13}$C MR imaging in diamond microparticles. We demonstrate high-fidelity imaging in either mode and show they can be rendered background-free. We propose three imaging regimes wherein such dual-mode imaging provides benefits over either mode considered individually. In particular, in either scattering or scattering-free media, optics and MRI can complement each other with respect to resolution, signal-to-noise ratio (SNR), and depth of imaging. We propose a hybrid sampling strategy, wherein conjugate imaging is carried out simultaneously in real space and k-space to enable image acceleration and power reduction in wide field-of-view settings.

Combined optical-MR imaging is made possible by special features of the diamond material medium. The diamond particles are incorporated with $\geq 1$ ppm of nitrogen vacancy (NV) defect centers (6). Under subbandgap illumination $<575$ nm, the particles fluoresce brightly in the red with high luminosity ($\sim 90$ cd/m$^2$) and optical stability. Fluorescence occurs concurrently with the optical polarization ($>10\%$) of the electronic spins.

Significance

We report on progress toward combining magnetic resonance imaging (MRI) and optical imaging in diamond microparticles. Our approach relies on the nitrogen vacancy (NV) center-driven optical nuclear hyperpolarizability in diamond particles, that renders them “bright” in MRI while simultaneously fluorescing optically. Both imaging modes allow suppression of background signals. We elucidate how such “dual-mode” imaging can perform better than either modes taken separately, leveraging relative strengths of optics and MRI with respect to resolution and imaging in scattering media. Finally, we propose a protocol for accelerated imaging that exploits the Fourier conjugacy in images obtained via optics and MRI. Our work suggests methods to combine optical and RF imaging in classes of deployable nanoparticles.
This can be transferred to $^{13}$C nuclei in the surrounding lattice, hyperpolarizing them and making them amenable to direct MR imaging (7). We exploit a recent mediated hyperpolarization technique (8) that allows large ($\sim$1%) $^{13}$C polarization levels in diamond particles at room temperature and low magnetic fields.

Results

Fig. 1A is a schematic of the experiment. Diamond particles (200 µm size, $\sim$40 mg) arranged in a ring-shaped phantom (Fig. 1D) are imaged optically under continuous 520-nm illumination and 630-nm long-pass filtering (Fig. 1E). The particles fluoresce brightly ($\sim$12 $\times$ 10$^{12}$ cps) (SI Appendix). The same optical excitation polarizes (initializes to $m_e = 0$) the electron spins, and microwave (MW) sweeps across the electron spin resonance (ESR) spectrum drive Landau–Zener (LZ) dynamics that transfer polarization to the $^{13}$C nuclei in an orientation-independent manner (Fig. 1B) (8, 9). We obtain $\sim$0.3% $^{13}$C spin polarization in 40 s under 1 W total optical illumination, enabling hyperpolarized MR imaging (Fig. 1B). This level corresponds to a signal enhancement of 280 times over thermal $^{13}$C polarization at 7 T (Fig. 1C) or 206 times over 9.4 T ($\sim$5 $\times$ 10$^4$ over 38 mT) and $\sim$10$^4$ acceleration in MR imaging time.

Compared to conventional dynamic nuclear polarization (DNP) techniques, our method requires relatively low laser ($\sim$2 mW/mg) and MW power ($\sim$0.05 mW/mg) (10). The MRI demonstration in Fig. 1F employed a laser power density of $\sim$80 mW/mm$^2$ to polarize $\sim$40 mg of diamonds. While this work is focused on diamond particle imaging, we note possible extension for in vivo studies assuming that specific absorption rate (SAR) can be controlled to safe limits, and the diamond particles can be eliminated from the body. In the context of potential in vivo applicability, we estimate a MW SAR of 1.1 $\times$ 10$^4$ W/kg [G]$^2$. Hyperpolarization efficiency scales approximately logarithmically with MW power beyond a particular threshold (8), indicating that the SAR can potentially be curtailed without severely degrading the DNP enhancement factors (SI Appendix).

We use a variant of fast low angle shot (FLASH) (11) to produce the MRI images at 9.4 T (Fig. 1B and F), with short echo times (0.5 ms) to accommodate short $T_2 \approx$1 ms of $^{13}$C in diamond. To eliminate pulse interference during the $\sim$200 µs gradient switching periods, imaging was performed without a slice selection gradient (Fig. 1B). The SNR of the MR image (Fig. 1F) is $\sim$4 in 16 scans (each scan preceded by hyperpolarization), limited by rapid $^{13}$C $T_2$ decay, low sample filling factor ($\approx$0.007), and laser-limited hyperpolarization. The use of dynamic decoupling sequences, such as quadratic echos (12), spin locking (13), or a recently developed approach to enhance free induction decay (FID) time exceeding 2 s (14), can improve the imaging SNR by at least an order of magnitude. If the application at hand permits higher optical powers close to saturation intensity, similar gains can be concurrently obtained. In particular,
materials advances would boost MR image SNR by an order of magnitude through $^{13}$C enrichment and through the use of high-temperature annealing (15). Further improvements may be realized by optimizing the detection coil geometry and filling factor, for instance through the use of small-volume inductively coupled receiver coils (16). These concerted gains in MR signal could also permit similar high-contrast images in nanosized particles ($\leq 100$ nm), although these smaller particles display lower ($\approx 10^{-2}$) hyperpolarizability than the microparticles employed in this work (15). The MRI spatial resolution here is 640 $\mu$m $\times$ 640 $\mu$m $\times$ 1 mm. The DNP method here presents advantages over traditional hyperpolarization methods for solids imaging, employed for instance in $^{29}$Si microparticles (17). We work at room temperature and low field ($\approx 40$ mT) and polarize samples in under 1 min of laser pumping. Conventional methods, in contrast, require high magnetic field ($\geq 3$ T) and low temperature ($< 4$ K) and polarization buildup can take several hours (18). While the absolute polarization is lower in our method, we circumvent the high polarization loss (as large as 99%) (19) accrued upon thawing and sample transfer out of the cryostat. Technologically, our technique aids end-user operation—MW amplifiers and sweep sources are low cost and readily available, and hyperpolarized particles can be delivered by a portable device (10). Since the DNP process is detection-field agnostic, the technique is especially interesting in the context of low-field MRI where hyperpolarization can be replenished continuously.

**Background-Free Imaging**

Both optics and MR modalities allow on-demand image amplitude modulation, enabling common-mode suppression of background signals. The NV fluorescence is conditioned strongly on the misalignment angle $\theta$ of the N-to-V axis to the applied field (20, 21), especially at low fields approaching 50 mT. This arises from mixing of the $m_s=\pm 1$ spin levels in the excited state. Since the randomly oriented particles sample all possible $\theta$ angles, this enables a simple method to modulate the optical images by applying a pulsed field $B_{ext}$ (22). In Fig. 2 A, $i$, we simulate fluorescence dependence under $B_{ext}$ using a seven-level model of the NV center (20): $\frac{dn_l}{dt} = \sum_{j=1}^{3} (k_{ij} n_j - k_{ji} n_i)$, where $n_i$ is the population of the $|i\rangle$ state, and $k_{ij}$ denotes the kinetic transition rate between states $|i\rangle$ and $|j\rangle$ (measured in ref. 20). In steady state, photoluminescence (purple line in Fig. 2 A, $i$), obtained evaluating $\int n_i(B_{ext}, \theta) \sin \theta d\theta$ decreases with $B_{ext}$, in reasonable agreement with (normalized) measurements (blue dots). In our experiments (Fig. 2B), $B_{ext}$ = 40 ± 2 mT takes the value identical to that used for hyperpolarization, and we obtain an $\approx 10\%$ optical modulation contrast (Fig. 2A, ii).

MRI mode allows for similar signal modulation. Such modulation refers to controlling the $^{13}$C hyperpolarization sign based on the direction of MW sweeps between alternate scans (as opposed to continuously modulated signal in optics). This effect originates from LZ dynamics excited by the chirped microwaves (Fig. 2C, 8, 9). The $^{13}$C nuclei are aligned (antialigned) with the polarization field under low-to-high (high-to-low) frequency sweeps. This allows complete sign reversal of the MRI images at full contrast. As a figure of merit, we characterize modulation contrast as the difference ratio of the MR images $I$ and $\bar{I}$ under opposite MW sweeps (Fig. 2D) as $\Delta = \frac{I - \bar{I}}{I + \bar{I}}$, where $N^2$ is the total number of pixels. From the data in Fig. 2E and F, we obtain $\Delta = (194 \pm 3)\%$. Similar modulation contrasts are challenging to achieve in conventional cryogenic DNP due to technical limitations of MW cavity switching (19, 23).

Signal modulation allows background-free imaging of the diamond particles. We refer to “background” in this context as media with fluorescence or $^{13}$C NMR signals that overlap in

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**Fig. 2.** On-demand dual-mode image modulation. (A, $i$) Normalized fluorescence signal for a randomly oriented diamond particle ensemble under an applied magnetic field (points, experiment; purple line, simulation). We ascribe the discrepancy to scattering effects. (A, ii) Optical modulation under 40 ± 2 mT pulsed magnetic field showing a signal contrast $\approx 10\%$. (B) Optical images under 0 and $\approx 40$ mT applied field showing weak $\approx 10\%$ optical contrast. (C) The $^{13}$C hyperpolarization sign control. MW frequency sweeps in low-to-high (high-to-low) fashion across the ESR spectrum lead to positive (negative) hyperpolarization. Shown are 7 T $^{13}$C NMR spectra under opposite sweep conditions. (D) The $^{13}$C MRI images under opposite MW sweep conditions, showing full sign reversal and $\approx 194\%$ modulation contrast. Here FLASH images were taken with TE = 0.6 ms and TR = 6 ms.

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wavelength (or NMR frequency) with the diamond particles. As a proof of concept (Fig. 3), we consider particles being concluded with a high concentration of Alexa 647, a fluorescent dye with emission in the 650- to 670-nm range (SI Appendix), as well as $^{13}$C-methanol, which has a chemical shift nearly overlapping that of diamond. These solution media fill both the inner and outer spaces of the capillary tube that compose the diamond phantom in Fig. 3 A and B. The backgrounds result in images that are circle shaped since the diamond phantom is completely indiscernible within it (Fig. 3 C and D). To recover the diamond signals in the optical mode, we perform lock-in detection under a 40-mT 0.1-Hz square-modulated field. We record a 2,000-frame movie at 0.1 fps and computationally apply lock-in suppression on each pixel. The resulting image (Fig. 3E) shows recovery of the diamond signal, in this case, from the 2 times stronger background. Optical background suppression is ultimately limited by the modulation contrast, and up to an order of magnitude is experimentally feasible (22). Concurrent MRI background suppression is realized by subtracting the images under opposite sweep-ramp hyperpolarization conditions in alternative scans. The $^{13}$C-methanol signal was 5 times larger than the diamond signal and is efficiently canceled (Fig. 3F).

**Regimes for Combined Dual-Mode Imaging**

While Fig. 3 shows the optical and MR imaging separately, what may one accomplish by combining them together in a dual-imaging modality? Consider that MRI is natively immune to optical scattering, while optics provide superior resolution to optical scattering-free media. This can make a combined modepersuasive in specific imaging regimes.

Consider first the fluorescent imaging of nanodiamond particles embedded in a scattering medium (e.g., in tissue). One has to contend with round-trip attenuation and scattering losses (scaling $\sim \lambda^{-4}$) that restrict the proportion of collected photons. In tissue, for instance, attenuation and scattering at 650 nm lead to exponential losses with coefficients 0.1 and 1.1 mm$^{-1}$, respectively (24, 25). We estimate a round-trip attenuation loss of 0.2 and scattering loss of $10^{-4}$ at 3 mm depth. There are further photon losses stemming from green–red conversion ($\sim 10^{-5}$) and the unusually high refractive index differential between diamond and its environment ($\sim 10^{-2}$), as well as the finite numerical aperture limited by geometric solid-angle constraints of the detector at high fields of view (FOVs) ($\sim 10^{-2}$) (SI Appendix). Scattering also leads to a simultaneous loss in imaging resolution. Random scattering of the beam blurs the optical image, spreading it as much as 800 Âµm at 1 cm depth (see SI Appendix for simulations).

In contrast, hyperpolarized MR imaging SNR can be competitively efficient, especially at increasing imaging depth $d$. One partakes of losses only in the one-way illumination of the particles with 520 nm light. Hyperpolarization efficiency is $\sim 2 \times 10^{-4}$ per $^{13}$C nucleus per incident photon (8). The high detection losses and geometric solid-angle collection constraints are replaced by more benign factors related to sample-coil filling, detector Q, and the overall MR detection frequency (26). Surface coils matched to the sample and the use of high-Q ferrite resonators can lead to substantially more efficient detection (27). In this regime MR imaging could have a higher overall SNR than its optical counterpart. This cross-over point in efficiency occurs at depths $d \approx \frac{1}{Q} \log(\eta_d)$ at 650 nm, where $\eta_d$ is the ratio of optical and MR imaging SNR for surface diamond particles. Immunity to scattering also means that MR imaging resolution is independent of imaging depth (SI Appendix).

To now elucidate advantages of dual-mode optical MR imaging, we consider the effect of one mode enhancing the other. We focus on two imaging regimes, conditioned on imaging in media with and without optical scattering, respectively. In each regime we consider a “primary” imaging mode (either optics or MRI) and wherein a “secondary” (complementary) mode is added to enhance it (Table 1).

**Regime 1.** Consider first optical imaging (primary mode) in scattering media. Scattering deteriorates image SNR by a factor

![Fig. 3. Dual-mode background suppression. (A and B) Schematic of imaging phantoms. Diamonds are arranged in a ring-shaped phantom, cosituated with Alexa 647 dye and $^{13}$C-methanol that present an artificial background for optical and MR imaging, respectively. (C and D) Optical and MR images with the background. Dashed lines serve as a guide to the eye for the imaging phantom. Diamond particles are indistinguishable from the background in both imaging dimensions. (E and F) Background suppressed optical and MR images employing signal modulation (reversal) allow complete recovery of the original diamond phantom in both imaging modes.](https://doi.org/10.1073/pnas.2023579118)
Table 1. Dual-mode imaging regimes from a combination of optics and MRI

| Particular specifics | Regime I | Regime II | Regime III |
|---------------------|----------|-----------|------------|
| Primary mode        | Optical imaging | Hyperpolarized MRI | Accelerated dual-mode imaging protocol |
| Secondary mode      | Hyperpolarized MRI | Optical imaging | |
| Operating regime    | Deep in scattering media | Shallow or in nonscattering media | Shallow or in nonscattering media |
| Primary mode SNR    | $\propto \gamma_0 \cdot e^{-\alpha_i d} e^{-\alpha_e d}$ | $\propto P_i (\Delta x_{\text{mri}})^2 \cdot \sqrt{T}$ | $\propto \sqrt{T/(1 - s)^{1/2}}^*$ |
| Dual-mode SNR       | $\propto e^{-\alpha_i d}$ | $\propto \frac{\text{FOV} \Delta x_{\text{optics}}}{\Delta x_{\text{mri}}} P_i (\Delta x_{\text{optics}})^2 \cdot \sqrt{T/\text{FOV} \Delta x_{\text{optics}}}$ | $\propto 1/(1 - s)^{1/4}$ |
| SNR gain            | $1/\gamma_0 \cdot e^{\alpha_i d}$ | $\frac{\text{FOV} \Delta x_{\text{optics}}}{\Delta x_{\text{mri}}}$ | $47.5 \sqrt{\text{Hz}}$ |
| Primary mode SNR (example) | $4.4 \times 10^{-3} \sqrt{\text{Hz}}^2$ | $0.63 \sqrt{\text{Hz}}^2$ | $47.5 \sqrt{\text{Hz}}^2$ |
| Dual-mode SNR (example) | $1.1 \times 10^{-2} \sqrt{\text{Hz}}^2$ | $6.3 \sqrt{\text{Hz}}^2$ | $47.5 \sqrt{\text{Hz}}^2$ |
| SNR gain (example)  | 2.5 | 10 | $5^*$ |
| Primary mode resolution $\delta x_p$ | $\propto d$ | $\delta x_{\text{mri}}$ | $\delta x_{\text{optics}}$ |
| Dual-mode resolution $\delta x_d$ | $\propto \frac{1}{\text{FOV} \Delta x_{\text{mri}}}$ | $\delta x_{\text{optics}}$ | |
| Resolution limit    | $\sim \frac{1}{\text{FOV} \Delta x_{\text{mri}}} (\sim 1 \mu m)^8$ | $\delta x_{\text{optics}}$ | N.A. |
| Primary mode resolution (example) | $1200 \mu m$ | $640 \mu m$ | N.A. |
| Dual-mode resolution (example) | $640 \mu m$ | $40 \mu m$ | $40 \mu m$ |
| Resolution gain (example) | 1.875 | 16 | N.A. |
| Power reduction $^\dagger$ | $1/\gamma_0 \cdot e^{\alpha_i d}$ | $\frac{\text{FOV} \Delta x_{\text{optics}}}{\Delta x_{\text{mri}}}$ | $\propto 1/(1 - s)^{1/2}$ |
| Power reduction (example) | 2.5 | 10 | $25^{**}$ |
| Background suppression | MRI can suppress optical background | Minimal background in $^{13}$C MRI | Field modulation can suppress optical background$^{11}$ |

Regimes I and II consider optics and MRI being primary imaging modes in scattering and scattering-free media, respectively. Regime III considers sampling in both imaging dimensions as elucidated in the algorithm of dual-mode imaging. Red color indicates parameters where image enhancements can be gained via dual-mode combination, while blue indicates less improvement. For clarity, the variables here refer to $\gamma_0$, ratio between optical SNR and MR SNR at $d = 0$; $P_i$, incident power density; $\Delta x$, pixel size; $\alpha_i$, $\alpha_e$, incident and emission light loss coefficients, including scattering and attenuation; $d$, object depth; $T$, imaging time; $\Delta x_{\text{optics}}$, $\Delta x_{\text{mri}}$, pixel size for optical and MR; FOV, field of view; and $s$, sparsity, the proportion of dark pixels.

$^*$ Extrapolated from SI Appendix, Fig. S2, and taking a depth of 15 mm.

$^1$ The SNR gain in the third mode is defined as the ratio between dual-mode SNR and optical SNR.

$^\dagger$ Extrapolated from experimental number to $d = 3$ mm, using loss coefficient 13.5 cm$^{-1}$ for 532-nm laser and 12.1 cm$^{-1}$ for 650-nm fluorescence in scattering media (see SI Appendix, section 1A, diamond mass $\sim 40$ mg). The coefficients are calculated based on fatty tissue data in ref. 32.

$^\ddagger$ Taken from the dual-mode experiment demonstrated in Fig. 1 (SI Appendix, section 1A).

$^\S$ Based on gyromagnetic ratio of $^{13}$C, $G_{\text{max}}$ can be up to 60 T/m in ref. 33, and $\tau \sim T_2 \approx 1$ ms.

$^\parallel$ Power reduction is defined as the ratio between the power required by the single primary mode and the dual-mode approach to achieve certain SNR.

$^\S^\S$ Obtained from Fig. 4 when assuming $(1 - s) = 0.9 \times 10^{-3}$.

$^\S^\S^\S$ See Fig. 3.

of $e^{-\alpha_i d} e^{-\alpha_e d}$, where $\alpha_i$, $\alpha_e$ are incident and fluorescent loss coefficients, respectively. Instead, the ability to image via the secondary MRI mode can improve SNR compared to the primary mode by a factor $e^{-\alpha_e d}$ and imaging resolution ($\propto d$). Table 1 shows these factors, with estimated numerical gains possible in the scenario with imaging depth $d = 3$ mm (see footnotes in Table 1). In fact, the depth penetrability of MRI and its relative immunity to scattering can also provide imaging capability in three dimensions.

**Regime II.** Alternately, consider when MRI is chosen as the primary mode and is applied in a scattering-free imaging regime. Here the resolution can be augmented from $\delta x_{\text{mri}}$ (640 $\mu m$ in Fig. 1) to $\delta x_{\text{optics}}$ (40 $\mu m$ in Fig. 1, ultimately limited by diffraction) by rastering a laser beam across the field of view to selectively hyperpolarize one single pixel at a time. Assuming the available total power is fixed, focusing the beam increases power density by a factor of $\frac{\text{FOV} \Delta x_{\text{optics}}}{\Delta x_{\text{mri}}}$. Despite pixel size change and imaging time extension, these factors can provide an SNR enhancement of $\frac{\text{FOV} \Delta x_{\text{optics}}}{\Delta x_{\text{mri}}}$ simultaneously with resolution gain.

**Accelerated Conjugate-Space Imaging**

While the discussion above considers dual-mode imaging in terms of relative merits of optics and MRI with respect to scattering and resolution, here we propose an additional regime (regime III) exploiting the native ability of the two imaging methods to sample in Fourier reciprocal spaces. This allows feedforwarding information from one space to guide the sampling in the other space. Every sampled point in one space carries

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information from all of the points in the other space; this is exploited to provide speedup in sparse imaging settings. This approach shares similarities with compressed sensing (28, 29) (see SI Appendix for a detailed comparison), but is different in the sense that image sampling here physically occurs in two conjugate spaces.

As a specific example, consider a wide FOV imaging scenario where optical imaging is performed by rastering a low-power beam across a sample of nanodiamonds distributed in a volume. We assume the objects are sparse in FOV and, for simplicity, that the per-pixel imaging time cost is identical for both optics and MRI. Fig. 4 describes the imaging protocol. Given the sparse original image to be acquired, a subset of \( \ell \) points in each dimension is first sampled in k-space via MRI. The resulting blurry k-space image is thresholded and fed forward to confine real-space points to be scanned over. The per-pixel imaging time cost is identical for both optics and MRI imaging in diamond particles. However, the time cost to be accrued per sample (pixel) is identical for both optical and MR imaging dimensions, although it is straightforward to scale the results by the cost ratio \( \eta \) as appropriate.

There is an optimal k-space sample threshold \( \ell_{opt} \), arising as a compromise between better constraining real-space sampling and taking longer to image. Fig. 4C demonstrates this for a 32 x 32-pixel FOV, where we consider normalized imaging time savings over either modality for target images with varying sparsity to identify \( \ell_{opt} \). We estimate that hybrid sampling can deliver more than an order of magnitude in time savings (right axis in Fig. 4C), while requiring only the scanning of \( \ell_{opt} \approx 10\% \) of total k-space (upper axis). As expected, \( \ell_{opt} \) decreases with sparsity, a reflection that larger k-samples are required with increasing image complexity (Fig. 4D). Given the small FOV and discrete values of k-samples, this manifests in the staircase-like pattern in Fig. 4D, but scales \( \ell_{opt} \approx (1-s)^{1/4} \) (solid line) as we shall derive below. Finally, in Fig. 4E we consider the combined imaging time \( \tau \) under optimized conditions as a function of image sparsity, assuming that time for optical imaging is 1. Indeed, the imaging acceleration can be quite substantial, scaling as \( \tau^{-1} \sim (1-s)^{-1/2} \).

We now elucidate origins of the imaging acceleration by studying the convergence trajectory of the reconstructed image as it approaches the target with each step of the protocol (Fig. 4F). Considering several image configurations with a fixed sparsity, we analyze in Fig. 4F the overlap of the reconstructed image to the target image through the correlation \( C = \sum (-\langle \cdot \rangle(-\langle \cdot \rangle)) \), where \( \langle \cdot \rangle \) indicates the mean value. Indeed, under usual rastered optical image acquisition (dashed green line in Fig. 4F), the reconstructed image linearly approaches the target as more
samples are acquired. In contrast, employing the hybrid acquisition of a few k-space points, and by constraining the spacing over which the final image is to be acquired, one obtains a rapid convergence with the target. Numerically, the slope of convergence scales approximately \( (1 - s)^{-1/2} \) (Fig. 4 F, Inset), indicating rapid gains can be amassed at high image sparsity. To analytically elucidate image acceleration at high sparsity (solid lines in Fig. 4 D and E), let us consider the real-space target image \( f(x, y) \), which in k-space is \( F[f] = \tilde{f}(k_x, k_y) \), where \( F \) denotes a spatial Fourier transform. As k-space sampling now occurs just to \( \ell \)th order, one obtains the reconstructed image, 
\[
\tilde{f}(k_x, k_y) = \tilde{f}_n(k_x, k_y) - \Pi \left( \frac{k_x}{s} \right) - \Pi \left( \frac{k_y}{s} \right),
\]
where \(\Pi \) is a rectangular function representing a sampling window with a side length of, for instance, \( W_{kx} = \delta k_x \ell \), where \(\delta k_x = 1/N_{kx} \) is the k-space pixel size. Transformation back to real space gives the convolution, 
\[
F^{-1}[\tilde{f}] = f(x, y) = \delta k_x \delta k_y s \left( W_{kx} \text{sinc}(W_{kx} x) - W_{ky} \text{sinc}(W_{ky} y) \right),
\]
an object of pixel radius \( r \) is effectively blurred in \( r \rightarrow r \frac{\sqrt{s}}{s} \), where the factor \( r_0 \) is set by the thresholding level employed (SI Appendix), and \( s/N \) is the effective k-space sampling ratio. Increasing \( \ell \) makes a more faithful representation and improves regional constraints, but it is associated with a time cost. To analytically evaluate the time savings let us as an example consider the FOV consists of \( n_{kx} \) objects of radius \( r \), giving \( (1 - s) = n_{kx} \pi r^2/N^2 \). The normalized imaging time is then \( \tau = (1 - s) + \frac{n_{ky}}{n_{kx}} \left( \left[ (1 - s)N \right]^2 + \frac{2\pi}{\delta k_x} \left[ (1 - s)N \right]^2 + \frac{2\pi}{\delta k_y} \right) \) (SI Appendix). In the limit of high sparsity, \( \tau \rightarrow 0 \), \( (1 - s)N \rightarrow 0 \), and \( (1 - s)N^2 \rightarrow O(1) \), giving \( \tau \approx C_0 \frac{1}{(1 - s)N^2} \), where \(C_0 = \frac{n_{ky}}{n_{kx}}\) is a constant. Determining the optimal \( \ell \) to minimize \( \tau \) gives \( \ell_{opt} \propto (1 - s)^{1/4} \), and the optimal (normalized) time \( \tau \propto (1 - s)^{1/2} \).

The scaling from this simple model is shown as the solid lines in Fig. 4 D and E and closely matches numerical results in the limit of high sparsity.

Finally, we comment that imaging acceleration results in a lower total optical power delivered to the sample by the same factor as the acceleration gain. This protocol might have real-world applications given that most imaging of targeted nanodiamond operates in the high-sparsity limit \( s > 95\% \) (30, 31).

Conclusions
We have demonstrated a method for dual-mode optical and MR imaging in diamond microparticles. Our approach relied on optically fluorescent centers that simultaneously spin polarize \(^{13}\text{C}\) nuclei, making the particles “bright” under MR imaging. We discussed means by which the two modalities can be combined, exploiting complementary advantages for scattering-free and high-resolution imaging. We finally proposed methods for image acceleration exploiting the Fourier conjugacy inherent to optics and MRI.

Data Availability. All study data are included in this article and/or SI Appendix.

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