Nitrogen-vacancy (NV) quantum magnetometers offer exceptional sensitivity and long-term stability. However, their use to date in distributed sensing applications, including remote detection of ferrous metals, geophysics, and biosensing, is limited due to the need to combine optical, microwave (MW), and magnetic excitations into a single system. Existing approaches have yielded localized devices but not distributed capabilities. In this study, a continuous system-in-a-fiber architecture is reported, which enables distributed magnetic sensing over extended lengths. Key to this realization is a thermally drawn fiber that has hundreds of embedded photodiodes connected in parallel and a hollow optical waveguide that contains a fluid with NV diamonds. This fiber is placed in a larger coaxial cable to deliver the required MW excitation. This distributed quantum sensor is realized in a water-immersible 90-m-long cable with 102 detection sites. A sensitivity of $63 \pm 5 \text{ nT Hz}^{-1/2}$ per site, limited by laser shot noise, is established along a 90 m test section. This fiber architecture opens new possibilities as a robust and scalable platform for distributed quantum sensing technologies.

Current distributed fiber sensors have high sensitivity to temperature, strain, and pressure but not to magnetic fields. Here, we add distributed spin-based quantum magnetometry to the sensing capabilities of distributed fiber sensors through the integration of NV ensembles. Recent years have seen the rapid advancement of NV solid-state quantum sensors because of their excellent sensitivity to magnetic fields, with sub-nT Hz$^{-1/2}$ sensitivity in the dc limit, high dynamic range vector resolution, and remarkable long-term stability. Recent advances toward component integration have achieved distributed magnetic point probes consisting of micro-diamonds on fiber facets or nanodiamonds on tapered fibers. However, practical devices will require compact and stable architectures and—for some applications—distributed sensor arrays for magnetometry. Here, we introduce an NV-based fiber magnetometer composed of hollow-core silica fiber, drawn with a functional cladding containing the necessary electronics for optical detected magnetic resonance (ODMR) magnetometry. We likewise use the fiber as a delivery mechanism for the pump field, allowing spin polarization and readout, while extending component integration by having the fiber serve as a backbone for photodetection and microwave spin control. The culmination of these properties within our fiber enables distributed magnetometry.

The device fabrication is outlined in Figure 1a. The process consists of two thermal fiber drawings. First, we draw a hollow-core silica fiber (index 1.46) through a high-temperature draw tower (2050 °C), which is subsequently polymer-coated (MyPolymer OF-145N, UV-curable index 1.445). The polymer coating functions as cladding for light guidance and mechanically protects the brittle silica shell. Next, we produce a polycarbonate preform with embedded, prefabricated photodiodes (PD) located in prescribed locations along the preform. During the low-temperature thermal draw at 280 °C, we unspool the hollow-core fiber together with two copper wires into separate prefabricated channels in the preform (see Figure 1a). As the preform is thermally drawn, the PDs separate axially while the two wires approach each other until making contact with the PDs. The PDs operate in a parallel circuit. We insert the finished fiber into a notched coaxial cable (RG6/U MOHAWK M71003), which supplies the driving MW fields to all the sensing sites simultane-ously.

By feeding the PC preform at 1 mm min$^{-1}$ while drawing at 0.8 m min$^{-1}$, we produce the completed fiber with the desired cross section of 0.9 × 0.9 mm. Figure 1b shows a close-up of the fiber with one of the PDs visible. The cross section in Figure 1c shows the fiber and copper wires, which straddle the PD (not visible in this cut) as indicated by the dashed lines. Figure 1d shows the 300 m fiber spool. The complete cable assembly is shown in Figure 1e.

The magnetometry relies on an ensemble of micro-diamonds containing NVs in an oil droplet (Nikon 50 Immersion Oil with an index of 1.518) that can be spatially scanned through the hollow-core fiber. We apply an oleophobic coating (3M Novec 1720 Electronic Grade Coating) to prevent adhesion of the micro-diamond-oil droplet to the inner silica wall. The 100-µm inner diameter of the silica fiber allows free movement of the oil drop;
the 200-μm outer diameter is chosen to enhance mechanical robustness.

This integrated fiber detection architecture spans decades of dimensional scales and different materials to deliver a distributed magnetometer system. Specifically, this fiber ranges from nanoscale NV centers hosted within micrometer diamond crystals, with 670 μm silicon PDs, to 100-m-scale cable. A cross section of the completed device is illustrated in Figure 2a. The fiber contains periodically spaced sensing modules (average spacing of 17 cm) to measure the magnetic field magnitude |B(s)| at distance s along the fiber. The 532-nm excitation field required for optical detection of magnetic resonances (ODMR) is guided in the silica shell (see Figure 2b).

When the waveguided pump reaches the high-index oil droplet, it is scattered into the micro-diamond-oil droplet, whose 1-mm length matches that of the embedded PD (Si PD T330P Vishay, and InGaAs/InP PD-TCT13 Three Five Materials). The integrated PDs differ from the chalcogenide photoresistive materials which have previously been drawn in fibers by having higher responsivity and bandwidth.[26,27] We scan the NV-micro-diamond droplet with air pressure supplied at the back end of the fiber. This droplet contains several thousand high-fluorescence micro-diamonds (Adamas). Figure 2c is a photograph of the NV droplet in the fiber. When the droplet is aligned with one of the embedded PDs, the red NV fluorescence is detected from the green pump background using a lock-in technique described below. The PDs are connected in parallel through two fiber-embedded copper wires, using a recently developed self-aligning fiber drawing process.[25] The surrounding coaxial cable drives the NV spin resonance for ODMR. The embedded PDs collect the emitted NV fluorescence, and the generated photocurrent is measured using a centralized readout bus located at the end of the fiber.

A key component of NV spin-based magnetometry is the spin dependence of the NV’s fluorescence. Figure 3a shows the NV spin triplet fine structure, which is the origin of the NV spin-dependent fluorescence. The frequency difference between |m_s = 0⟩ and |m_s = ±1⟩ magnetic sublevels is given by Δ_m = ± 2γB_z, where Δ_m = 2.87 GHz is the NV zero-field splitting, γ ≈ 28 GHz T−1 is the NV gyromagnetic ratio, and B_z is the applied magnetic field along the NV axis. The NV fluorescence intensity depends on its spin: it is “bright” in the |m_s = 0⟩ state due to radiative cycling transition and “dark” in |m_s = ±1⟩ due to channeling into a metastable singlet state, which ultimately decays into the |m_s = 0⟩ sublevel. A microwave field moves the ground state population between the |m_s = 0⟩ and |m_s = ±1⟩ levels when resonant. This ODMR spectrum[28,29] is shown for a single micro-diamond with zero magnetic field in Figure 3b. A resonance is observed at Δ_m; the contrast, C, represents the change in normalized fluorescence intensity with (black) and without (grey) applied MW field.
To determine the magnetic field, we modulate (at 6 kHz, 100% depth) the amplitude of the MW field and monitor the NV fluorescence using a lock-in amplifier. The measured lock-in signal is proportional to $C$.

The micro-diamond ensemble (photographed in Figure 3d-inset) within our droplet has multiple NV orientations. Each NV orientation experiences different $B_z$ and thus has different $|m_s| = \pm 1$ resonant frequencies. This diversity of resonant frequencies causes $C$ to decrease as a function of $|B|$ and be insensitive to the direction of $B$.[30] The NV fluorescence gave a current of about 5 nA for an ambient magnetic field which was fed through a 10 MOhm transimpedance amplifier (TIA). Figure 3c shows the lock-in signal as a function of magnetic field. Monitoring the lock-in signal at $D_0$ (dashed black line) gives a mapping between the lock-in signal and $|B|$ (see Figure 3d).

We use the mapping in Figure 3d to measure a 40 s period square wave applied with an external electromagnet (see Figure 3e). The measured noise is 55 nT, which extrapolates to a sensitivity of $63 \pm 5$ nT Hz$^{-1/2}$ when accounting for the 100 ms time constant and the 0.78 Hz equivalent noise bandwidth of the lock-in amplifier (24 dB/oct roll-off, see Supporting Information). The PDs other than the one performing the sensing are not exposed to either the guided light nor ambient light as the coaxial cable is opaque. The dark current from all the parallel-connected PDs is below the shot noise from the sensing PD.

To showcase the distributed quantum magnetometry capabilities of our device, we deploy it down a 70 m corridor (Figure 3f). We measured the magnetic field at 102 points along the corridor. We compare our measurement to a hand-held magnetometer (AlphaLabs Gaussmeter Model VGM) as seen in Figure 3g. We observe a spike in the magnetic field of $\approx 180 \mu$T at the location of Airgas cylinders. Figure 3g-inset shows the measured MW loss as a function of MW frequency for the cable assembly. We measure a loss of 49.5 dB over 90 m, which extrapolates to $=0.5$ dB m$^{-1}$ at 3 GHz. In principle, the MW loss could be further reduced by using better coaxial cable at the expense of the diameter of the cable system.

Our fiber-integrated magnetometer is sealed within a waterproof cladding, making it resilient under adverse conditions. Figure 4a is a photograph of the cable assembly placed into a water tank. A magnet was placed in the water and moved horizontally using a micrometer. The lock-in signal as a function of MW frequency for different magnetic fields is shown in Figure 4b. After 3 days of measurements within the water tank, we did not notice a degradation in the fiber magnetometry performance (see Figure 4c). We notice that moving the droplet in and out from the PD location and performing repeated measurements yields a systematic error that equals the sensitivity of our magnetometer.

Finally, we use the unique distributed nature of our fiber magnetometer to localize and quantify a magnetic source in space. Figure 5a shows the experimental schematic where a magnet cube ($3 \times 3 \times 3$ mm) is placed 1.3 cm away from the fiber. To demonstrate the capability to localize a magnetic object spatially, we align the fiber so that adjacent sensing modules are $3, 2.2, 5.7$, and $3$ cm apart, respectively, by making loops in the fiber. Figure 5b shows the measured lock-in signal as a function of MW frequency at each of the sensing modules. The fit (grey) of the spectra determines the applied magnetic field at...
Figure 3. ODMR spectra and level structure. a) Energy level diagram of the NV center. Magnetic fields are determined by measuring the NV resonance frequency for the $|m_s = \pm 1\rangle$ states with radiative (solid lines) and nonradiative (dashed lines) transitions shown. b) Measured ODMR spectrum of a micro-diamond with (black) and without (grey) MW fields applied taken using a microscope. C represents the change in measured normalized intensity due to the application of MW fields. c) Lock-in output of NV micro-diamond ensemble as a function of magnetic field, determined by a fit to the theoretical model (fit not shown). The left inset is a microscopy image of a single micro-diamond. d) We place an electromagnet near the cable and we measure the lock-in signal from the fiber magnetometer while changing the electromagnet magnetic field, calibrated in the inset. d-inset) We sweep the electromagnet current while measuring the magnetic field. e) We measure the magnetic field applied by an external electromagnet by monitoring the lock-in signal at $D_{21}$ (dashed line in c). The applied magnetic field is a 40 s period square wave with an amplitude of $\approx 60 \mu T$. The measured standard deviation is 55 nT. The histogram of the deviations from the mean is shown in the inset. f) We deploy the cable assembly in a 70 m corridor, and g) measure the magnetic field magnitude at every meter. We compare the quantum fiber magnetometry results (black dots) to a hand-held magnetometer (red error bars). The quantum fiber magnetometry error bars are smaller than the point’s shown. We measure a peak in the magnetic field at the location of the gas cylinders. g-inset) We measure the MW loss of the notched coax (black) and compare it to the loss of the untreated cable as reported by the manufacturer (red).

Figure 4. Underwater magnetometry. a) Photograph of the experiment. We insert our cable magnetometer into a water tank where its waterproof cladding protects it. Inset—zoom in on the notched cable where the PD is visible, and a diamond droplet is located. b) Lock-in signal as a function of MW frequency at different magnet distances. By moving the magnet horizontally, we change the magnetic field distance. The color represents the magnetic field values determined from the ODMR curves. c) We remove the magnet and perform repeated measurements over 3 days.
each node, giving the magnetic field distribution along the fiber. Modeling the applied magnetic field as a magnetic dipole, allows us to determine the location, orientation, and magnitude of the magnetic object (see Supporting Information). For example, we determine the axial and radial position with an accuracy of 1.87 and 2.87 mm, respectively. We compare the numeric solution (red) to direct measurement (black) in Figure 5c. The magnetic field magnitude along the fiber magnetometer is relatively insensitive to the magnet orientation; therefore, we observe error bars of $\pm 10^\circ$ for the orientation parameters $\theta$ and $\beta$.

Moving the droplet axially within the fiber in a reversible manner is done at typical speeds between 10 and 30 mm s$^{-1}$ due to the hollow-silica dimensions and the oil viscosity, which is comparable to similar microfluidic optical fibers. The scanning time can be decreased by increasing the diameter of the fiber at the cost of collection efficiency, as the diode droplet distance increases. Another approach to improve the scanning speed is by incorporating multiple droplets within the hollow fiber, where only one droplet is aligned with a sensing site. The average spacing between photodiodes is 17 cm, allowing up to 170 droplets simultaneously within our device for a droplet size of 1 mm. In each measurement instance, only one droplet will be aligned with a photodiode. Moving the 170 droplets at a speed of $\approx 10$ mm s$^{-1}$ allows a total scanning time of less than 10 min across a 1 km cable. However, there is a tradeoff between scanning speed and sensitivity: when working with multiple droplets, it is required to decrease their size and limit their amount due to optical losses from strong scattering in each droplet location (we measure a loss of 2.4 dB per 1 mm droplet); this means the sensitivity will be degraded due to optical loss. To achieve simultaneous sensing from multiple detection sites using the presented technique, we propose to modify the PD chip and to include a modulator in them. By that, each PD will transmit its measurement in a different modulation frequency, and the
separation can be made using a lock-in. Further improvements can be made by integrating custom CMOS-NV quantum sensing chips for localized spin control and readout.\[32\]

Several improvements in the sensitivity of this fiber magnetometer can be made. As shown in Figure S3, Supporting Information, the dominant noise source in our experiment is shot noise from the pump laser. We estimate that an additional order of magnitude improvement in sensitivity is possible by improving green filtering by \(\approx 20\) dB. Two possible solutions to filter the green light are inserting a thin free-space optical filter within the cladding or modifying the cladding to have a phononic Bragg grating layer.\[33\] The noise would then be dominated by the shot noise from the NV centers (see the purple line in Figure S3, Supporting Information). The SNR of this sensor could also be improved by increasing the total amount of collected NV fluorescence as shown in bulk diamond NV ensemble magnetometers, which have recently demonstrated a sensitivity of 290 pT Hz\(^{-1/2}\) for DC magnetometry.\[34\] One can achieve additional improvement by utilizing NV centers with improved spin coherence. These NVs would have steeper slopes than the one shown in Figure 3c, at the expense of dynamic range. Such NV centers are found in high-quality nanodiamonds,\[35,36\] within a CVD bulk diamond,\[37\] or an isotopically purified diamond.\[38\]

Several attempts have been made to avoid the need for microwave fields in NV-based magnetometry.\[39–41\] However, these techniques require both large bias magnetic field at each node and the use of single crystal diamond where the NV’s axis is aligned with the bias field. Other all-optical approaches such as coherent population trapping\[42\] or electromagnetically induced transparency\[43\] appear to be only compatible with a cryogenic operation. For those reasons, we chose to integrate the MW transmission line in the shape of coaxial cable throughout our fiber.

In conclusion, we have demonstrated a distributed, fully integrated magnetic field sensor that enables localization and quantification of magnetic fields along a 100 m scale length with a sensitivity of 63 nT Hz\(^{-1/2}\) and a spatial resolution of 17 cm. Such a distributed water-immersible quantum fiber magnetometer promises new applications for remote detection and tracking in a range of fields including geophysics, ferrous metal detection, and biomedical sensing.

Supporting Information
Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest
The authors declare no conflict of interest.

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