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Published in:
Physical Review Research

DOI:
10.1103/PhysRevResearch.1.033087

Publication date:
2019

Document version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
Detection of low-conductivity objects using eddy current measurements with an optical magnetometer

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Detection and imaging of an electrically conductive object at a distance can be achieved by inducing eddy currents in it and measuring the associated magnetic field. We have detected low-conductivity objects with an optical magnetometer based on room-temperature cesium atomic vapor and a noise-canceling differential technique which increased the signal-to-noise ratio (SNR) by more than three orders of magnitude. We detected small containers with a few mL of salt water with conductivity ranging from 4–24 S/m with a good SNR. This demonstrates that our optical magnetometer should be capable of detecting objects with conductivity <1 S/m with a SNR >1 and opens up new avenues for using optical magnetometers to image low-conductivity biological tissue including the human heart which would enable noninvasive diagnostics of heart diseases.

DOI: 10.1103/PhysRevResearch.1.033087

Optical magnetometers [1,2] based on laser interrogation of cesium or rubidium vapor can detect magnetic fields with sub-fT/√Hz sensitivity [3–6]. This high sensitivity is particularly useful for biomedical applications where tiny magnetic fields from the human body are detected. For example, optical magnetometers have detected brain activity [7–9], the heartbeat from adults [10] and fetuses [11,12], and nerve impulses [13]. Optical magnetometers can potentially also be used to noninvasively image the electrical conductivity $\sigma$ of the heart [14] using a technique called magnetic induction tomography (MIT) [15,16]. In MIT of the heart, one or more coils are used to induce eddy currents in the heart and an image of the heart is constructed from measurements of the associated induced magnetic field. This is a challenging task for several reasons, with the main one being the low conductivity $\sigma \lesssim 1 \text{ S/m}$ of the heart [14].

Imaging of low-conductivity objects has previously been done using coils for inducing and detecting the eddy currents. Large containers ($\approx 500 \text{ mL}$) with salt water with conductivity as low as 0.7 S/m have been imaged [15,17,18], and more recently, the spinal column has been imaged with a single scanning coil [19]. Optical magnetometers have several advantages compared to induction coils, in particular, they are widely tunable and can achieve high sensitivity which is fundamentally independent of the operating frequency. This is in contrast to induction coils which are sensitive to the change in magnetic flux and therefore have worse sensitivity the lower the frequency. For frequencies below 50 MHz, an optical magnetometer should be fundamentally more sensitive than an induction coil of similar size [20]. So far, optical magnetometers have been used to image highly conductive metallic samples ($\sigma \approx 10^6–10^8 \text{ S/m}$) [21–23] and also recently semiconductor materials with conductivities in the $\sigma = 50–10\,000 \text{ S/m}$ range [24].

In this work, we introduce a differential technique which improves the signal-to-noise ratio (SNR) by more than three orders of magnitude and then demonstrate detection of small containers with 8 mL of salt water with conductivity as low as 4 S/m. This represents a significant improvement compared to previous results with optical magnetometers and is a big step toward magnetic induction tomography of biological tissue with optical magnetometers.

We first discuss the standard approach for detecting and imaging a conductive object, in our case a container with salt water. Later we will discuss the differential technique. Consider a conductive object, a magnetometer, and a coil [denoted coil 1 in Fig. 1(a)] that generates a primary magnetic field $B_1(r,t)$ oscillating at the frequency $\omega = 2\pi \nu$. The primary field induces eddy currents in the object which in turn generate a secondary magnetic field $B_{es}(r,t)$. One can measure the total field $B(r,t) = B_1(r,t) + B_{es}(r,t)$ and by scanning the magnetometer or the object around it is possible to construct an image of the conductivity [21–23]. Varying the frequency $\omega$ can be useful for three-dimensional (3D) imaging [14] and for material characterization [22].

It is instructive to note that the primary field is attenuated while penetrating into the object due to the skin effect. The skin depth is $\delta(\omega) \approx \sqrt{2\pi / \mu_0 \omega \sigma}$, where $\mu_0$ is the vacuum permeability and we assumed that the object is nonmagnetic. When the thickness $t$ of the object is much smaller than the skin depth $\delta(\omega)$, the secondary field is 90° out of phase with the primary field and the ratio $\alpha$ of the amplitude $B_{es}(r_0)$...
of the secondary field to the amplitude of the primary field $B_1(r_0)$ at the magnetometer position $r_0$ is [15,25]

$$\alpha \equiv B_{xc}(r_0)/B_1(r_0) \approx -\Delta \sigma \omega \mu_0 \approx -2A/|\beta|, \quad (1)$$

where $A$ is a geometrical factor with dimensions of length squared. For a (2 cm)$^3$ container with salt water with conductivity $\sigma = 10.7 \text{ S/m}$ we calculate $\delta = 11 \text{ cm}$ and estimate $|\alpha| \approx 1.5 \times 10^{-4}$ when the frequency is 2 MHz (see Appendix). We demonstrate that it is possible to detect such a small change in signal with an optical magnetometer when using a differential technique. A more detailed calculation of the skin depth as a function of frequency for the case of a (2 cm)$^3$ container with salt water is given in Ref. [14].

The key component of our magnetometer is a paraffin-coated cesium vapor cell with a (5 mm)$^3$ inner volume [26]. The cesium atoms are spin polarized in the $x$ direction using circularly polarized pump and repump light and are detected using linearly polarized probe light [see Figs. 1(b) and 1(c)]. We denote the total angular momentum in the $F = 4$ hyperfine ground-state manifold $J$ and full polarization corresponds to $J = J_{\text{max}} = 4N_A\hat{x}$, where $N_A$ is the number of cesium atoms. The atoms are placed in a static magnetic field $B_0 \hat{z}$ and we are interested in detecting an oscillating magnetic field $B_1(t) = [B_1 \cos(\omega t) + B_1 \sin(\omega t)] \hat{y}$. The time evolution of the atomic spins is modeled using the Bloch equations with additional pumping and relaxation terms [27]

$$\frac{dJ}{dt} = \gamma J \times B + \Gamma_p J_{\text{max}} - (\Gamma_p + \Gamma_pr + \Gamma_{\text{dark}}) J, \quad (2)$$

where $\gamma$ is the cesium gyromagnetic ratio, $B = B_0 \hat{z} + B_1(t)$, $\Gamma_p$ is the rate of optical pumping, $\Gamma_{\text{dark}}$ is the decay rate in the absence of light, and $\Gamma_pr$ is the decay rate due to the probe light. We solve the differential equation in the frame rotating around the $x$ axis at the frequency $\omega$. Denoting the spin vector in the rotating frame $J'$ and assuming a steady state $\frac{dJ}{dt} = 0$, we find the spin components

$$J_{ss}' = J_{ss} \frac{\Delta^2 + (\delta \omega)^2}{\Delta^2 + (\delta \omega)^2 + \gamma^2(B_s^2 + B_c^2)/4}, \quad (3)$$

$$J_{ss}' = -J_{ss} \frac{\gamma(B_s \Delta + B_c \delta \omega)/2}{\Delta^2 + (\delta \omega)^2 + \gamma^2(B_s^2 + B_c^2)/4}, \quad (4)$$

$$J_{ss}' = -J_{ss} \frac{\gamma(B_s \Delta - B_c \delta \omega)/2}{\Delta^2 + (\delta \omega)^2 + \gamma^2(B_s^2 + B_c^2)/4}. \quad (5)$$

Here, $\delta \omega = \Gamma_p + \Gamma_{pr} + \Gamma_{\text{dark}}$, $\Delta = \omega - \omega_L$ is the detuning of the applied frequency from the Larmor frequency $\omega_L = B_0$, and $J_{ss} = J_{\text{max}} \Gamma_p/(\Gamma_p + \Gamma_{pr} + \Gamma_{\text{dark}})$.

If we only consider $B_c$, (i.e., $B_p = 0$), we see that $J_{ss}'$ and $J_{ss}''$ have dispersive and Lorentzian line shapes, respectively, as a function of detuning. The total width of the resonance is $\delta \omega = 1 + [B_1/B_{\text{sat}}]^2$ where $B_{\text{sat}} \equiv \Delta \omega/\gamma$. This means that the resonance is power broadened by the oscillating magnetic field $B_1$. If the magnetic field is on resonance ($\Delta = 0$), we have $J_{ss}' \propto \gamma_0 (1 + [B_1/B_{\text{sat}}]^2)$ which means that $J_{ss}'$ is only linear with the magnetic field for small fields $|B_1| < B_{\text{sat}}$.

The atoms are probed with linearly polarized light which due to the Faraday effect is rotated by an amount proportional to the spin component along the probe propagation direction. The light polarization rotation is measured with a balanced detection scheme leading to the magnetometer signal

$$S(t) \propto J_x(t) = \sin(\omega t) J_x'(t) + \cos(\omega t) J_y'(t). \quad (6)$$

The rotating spin components $J_x'$ and $J_y'$ are extracted from the magnetometer signal using lock-in detection at the frequency $\omega$. The lock-in provides an in-phase output $X \propto J_x'$ and an out-of-phase output $Y \propto J_y'$.

We characterize the magnetometer (without any conductive object) by applying the magnetic field $B_1(t) = B_1(r_0, t) = B_1(r_0) \cos(\omega t)$ Figure 2(a) shows the lock-in outputs as a function of frequency. The $X$ and $Y$ outputs have Lorentzian and dispersive line shapes centered around the Larmor frequency $\nu_L \approx 1.978 \text{ kHz}$ as expected from Eqs. (4) and (5) when $B_c = B_1(r_0)$ and $B_0 = 0$. The small side resonances (toward lower frequencies) are due to the nonlinear Zeeman effect [28,29]. The data set labeled “$B_1$” in Fig. 2(b) shows the lock-in outputs when the oscillating magnetic field $B_1$ is on resonance ($\Delta = 0$). We see that the mean values are $\langle X \rangle = 1.33 \text{ V}$ and $\langle Y \rangle \approx 0$ and that there is a significant amount of noise in the $Y$ output. In order to characterize the noise, we calculate the Allan deviation [30] of the $Y$ output which is roughly independent of averaging time with the value $\Delta Y_{\text{Allan}} = 22 \text{ mV}$ [see Fig. 2(c)]. The noise is mainly due to temporal fluctuations in the $B_0$ field. A change $\Delta B_0$ in the $B_0$ field shifts the Larmor frequency which then changes the $Y$ output. Close to the Larmor frequency $Y \approx a(v - \nu_L)$ where $a = -7.8 \text{ V/kHz}$ [see Fig. 2(a)] which means that a small change of the $Y$ output of 22 mV corresponds to a shift in the Larmor frequency of 2.8 Hz and a relative change in the $B_0$ field of $\Delta B_0/B_0 = 1.4 \times 10^{-6}$. This small number illustrates that an optical magnetometer requires a very stable $B_0$ field in order to precisely measure an oscillating magnetic field.
when the container is either empty or filled with salt water. The conductivity of the water can be conveniently varied between 0–24 S/m. All experiments were performed at a constant frequency of 1 kHz. Two measurements are shown in Figs. 1(a) and 1(b), with the container either empty or filled with salt water. In order to mitigate the above-mentioned problems, we introduce a differential technique where we use a second coil (denoted coil 2 in Fig. 1(a)) that generates a magnetic field \( B_2(\mathbf{r}_0, t) \) such that in the absence of the conductive object, the total magnetic field \( B_1(\mathbf{r}_0, t) + B_2(\mathbf{r}_0, t) \approx 0 \) at the position of the vapor cell. Coil 2 is placed further away from the conductive object than coil 1 such that eddy currents are mainly generated by coil 1 only. With this technique, the magnetometer signal is zero in the absence of the conductive object and the magnetometer should not be affected by power broadening or nonlinearities as long as the field from the eddy currents is smaller than \( B_{\text{sat}} \) [see Eqs. (4) and (5) with \( \Delta = 0 \), \( B_t = 0 \), and \( B = B_{\text{ec}}(\mathbf{r}_0) \)]. Furthermore, the signal-to-noise ratio of the measurement will improve by a factor \( 1/|\alpha| \) if the noise in the magnetometer is proportional to the total signal. This is a dramatic improvement as \( |\alpha| \approx 10^{-4} \) for our measurements on salt water.

Figure 2(b) shows three data sets. We see the noisy signal when \( B_1 \) is applied. When the opposite magnetic field \( B_2 \) is applied, the lock-in outputs change sign. Applying both magnetic fields \( 10(\mathbf{B}_1 + \mathbf{B}_2) \approx 0 \) at the same time (and increasing the amplitudes by a factor of 10) gives lock-in outputs close to zero with significantly reduced noise. Coils 1 and 2 are connected to two outputs of the same function generator and the amplitude and phase of the two outputs can be precisely set in order to zero the lock-in outputs. In Fig. 2(c) we see that the Allan deviation is \( \approx 130 \) times smaller for integration times \( \tau \geq 1 \) s when applying both magnetic fields compared to only applying \( B_1 \). Taking the factor of 10 into account, we find an improvement in signal-to-noise ratio of 1300 if detecting a low-conductivity object and a smallest detectable relative signal of \( |\alpha|_{\text{diff}} \approx 1.3 \times 10^{-5} \) when using the differential technique.

We now continue with detecting salt water inside a small container using our differential technique. The conductivity of the water can be conveniently varied between 0–24 S/m by changing the concentration of salt. Using a motorized translation stage, we scan the container 50 mm in the \( x \) direction a few mm above coil 1. Real-time traces of the \( Y \) output when the container is either empty or filled with salt water with varying conductivity are shown in Figs. 3(a)–3(d). With salt water present, we clearly see a change in the \( Y \) output when the container is on top of coil 1 (at the time around 10 s). The \( X \) outputs (not shown) are close to zero and do not change during the scan (within the statistical uncertainties). In order to reduce noise, the container is scanned \( \approx 20 \) times.
FIG. 3. Detection of salt water with the differential technique. (a)–(d) Real-time recordings. Each data point is integrated for \( \tau = 40 \) ms. (e) Relative change in signal using \( \approx 20 \) averages. Data are binned according to their position with one binned data point per 1 mm.

over coil 1 and the recorded traces are averaged. Figure 3(e) shows the relative change in signal \( \alpha \) for the averaged traces as a function of position. In order to guide the eye and to extract the maximum change in signal, we fit the data with salt water to a Gaussian function. For the \( \sigma = 10.7 \) S/m data we have the maximum change \( |\alpha| = 1.0 \times 10^{-4} \) which agrees reasonably well with the expected value (see Appendix).

We emphasize that we detect the salt water with good signal-to-noise ratio (SNR). We calculate the SNR as the maximum change in signal divided by the standard deviation (found from the data recorded with an empty container). For the traces in Figs. 3(b)–3(d) we have the SNR of 0.8, 2.5, and 6.1 for the conductivities 3.8, 10.7, and 24.1 S/m. For these traces, the integration time was only 40 ms. For the average traces in Fig. 3(e) we have the SNR of 6.4, 20, and 46. This demonstrates that our setup should be capable of detecting objects with conductivity \( \sigma < 1 \) S/m with a SNR >1 and that it should be possible to detect and image biological tissue which has conductivity \( \sigma \lesssim 1 \) S/m with our optical magnetometer.

The maximum change in signal as a function of conductivity is plotted in Fig. 4(a). We observe a linear dependence as expected from Eq. (1) confirming that the small observed signals are due to the salt water. We also vary the applied frequency (while at the same time adjusting the bias field to fulfill the resonance condition \( \omega = \gamma B_0 \)), and as shown in Fig. 4(b) we again observe the expected linear behavior. Finally, we vary the amplitude of the applied field. The signal starts out growing linearly but then some saturation occurs for higher amplitudes [see Fig. 4(c)]. The data are fitted to the function \( cU / (1 + [U/U_{sat}]^2) \) and we extract the saturation parameter \( U_{sat} = 5.2(2) \) V. This saturation is not expected when using the differential technique. We note that when only \( B_1 \) is applied, saturation happens at 10 times lower amplitudes. To avoid issues related to saturation, we used the amplitude \( U = 1 \) V for all other differential measurements (and \( U = 0.1 \) V for all measurements with one coil only).

In conclusion, we have demonstrated detection of small containers with salt water with conductivity ranging from 4–24 S/m using an optical magnetometer and a differential technique which improved the signal-to-noise ratio by more than three orders of magnitude. Our measurements were performed inside a magnetic shield, however, we expect that the differential technique will yield a larger improvement in unshielded conditions [31] as there will be more magnetic field noise which can be canceled. The technique also gives a large improvement when detecting objects with high conductivity (such as metal objects) as long as the detected field from the eddy currents is small compared to the primary field which for instance is the case when the object is far away. We note that similar techniques for canceling magnetic fields

FIG. 4. Detection of salt water with the differential technique. (a), (b) Relative change in signal as a function of conductivity and applied frequency. Data are shown together with linear fits. (c) Change in signal in mV as a function of the set amplitude \( U \) on the function generator connected to the two coils. Data are shown together with a fit to the function \( cU / (1 + [U/U_{sat}]^2) \) (solid line) and the linear part of the fit function \( cU \) (dashed line). A 1 V set amplitude corresponds to the fields 10\( B_1 = -10B_2 = 45 \) nT peak-to-peak amplitude.
have been used for imaging of structural defects in metal samples [32] and for detecting magnetic particles [33]. By further optimizing the sensitivity and long-term stability of our magnetometer, we expect that high-resolution imaging of biological tissue will be possible. This will make optical magnetometers promising candidates for localizing conduction disturbances in the heart, allowing for noninvasive diagnostics of heart diseases such as, for example, atrial fibrillation [14].

We would like to thank M. A. Skarsfeldt for preparing the salt-water solutions and B. H. Bentzen and S.-P. Olesen for motivating discussions. This work was supported by Innovation Fund Denmark through the Quibiz program, the ERC AdG QUANTUM-N, and the EU project MACQSIMAL.

APPENDIX

We now estimate the magnitude of the induced magnetic field generated by the eddy currents in the salt water using a semianalytical approach based on the calculations in Ref. [15]. The real container with salt water is cubic with (2 cm)³ inner volume. However, in the calculations below, we will for simplicity assume that the container is a cylinder with radius ρ and height t (also called thickness). The eddy currents are generated by the primary magnetic field \( \mathbf{B}_1 \) from coil 1 (see Fig. 5). The container and the coil are aligned on the same axis in the y direction. The coil has multiple windings, but we start by calculating the eddy currents generated by a single winding of radius \( r_w \) placed at a distance \( a \) from the container. Assume that an alternating current \( I(t) = I_0 \exp(i \omega t) \) with amplitude \( I_0 \) and frequency \( \omega \) is running through the winding. First, we calculate the induced eddy currents in a ring of radius \( \rho’ \) at a distance \( a + \tau’ \) from the winding, and with radial and axial thickness of \( d \rho’ \) and \( d \tau’ \), respectively. The induced eddy current is

\[
dI_{ec}(\rho’) = J d\rho’ d\tau’,
\]

where \( J \) is the current density. Note that \( J = \sigma E \), where \( \sigma \) is the conductivity and \( E \) is the magnitude of the electric field. The induced electromotive force \( E \) is first given by the integral of the electric field along a closed loop \( \oint \mathbf{E} \cdot d\mathbf{l} \). Second, it is given by \( E = -d\Phi/dt \), the time derivative of the magnetic flux \( \Phi = \int \mathbf{B}_1 \cdot d\mathbf{A} \) of the primary magnetic field \( \mathbf{B}_1 \) through the enclosed area. An analytical expression for the magnetic field \( \mathbf{B}_1 \) from the winding is given in Ref. [34]. With the assumption that this magnetic field changes instantaneously with the current in the coil, we find

\[
dI_{ec}(\rho’)
= -\frac{i \omega \sigma}{\rho’} \exp(i \omega t) \left( \int_0^{\rho’} B_1^{(y)}(a + \tau’, \rho’’) d\rho’’ \right) d\rho’ d\tau’,
\]

where \( B_1^{(y)}(a + \tau’, \rho’’) \) is the amplitude of the y component of the magnetic field from the winding at axial and radial distances \( a + \tau’ \) and \( \rho’’ \), respectively.

To calculate the magnetic field at the center of the vapor cell (distance \( b \) from the container) due to the eddy current in a ring of radius \( \rho’ \) we use the simple expression for the on-axis magnetic field from a current loop

\[
dB_{ec}^{(y)}(b + \tau’, 0) = \frac{\mu_0 \rho’^2 dI_{ec}(\rho’)}{2(\rho’^2 + (b + \tau’)^2)^{3/2}}.
\]

The magnetic field at the position \( r_0 \) of the center of the vapor cell from all the eddy currents in the whole salt-water sample is found by integrating the fields from the individual rings

\[
B_{ec}^{(y)}(r_0) = \int_0^b \int_0^\rho dB_{ec}^{(y)}(b + \tau’, 0)d\rho’ d\tau’.
\]

We then add the magnetic fields induced by the individual windings of the coil with their respective distances.

The model predicts that the induced field is shifted in phase by 90° with respect to the primary field, which agrees with our measurements. We numerically calculate the ratio of the amplitude of the induced magnetic field to the amplitude of the primary magnetic field taking all the windings of coil 1 into account. The calculation yields

\[
\left| B_{ec}^{(y)}(r_0) \right|/\left| B_1^{(y)}(r_0) \right| = 1.5 \times 10^{-4},
\]

using our experimental parameters (see Fig. 5), the frequency of 2 MHz, and the conductivity of 10.7 S/m. The calculated ratio is 50% higher than the experimentally obtained value. The main uncertainty on the calculated value stems from the distance between the container and coil 1. This distance could only be determined with an uncertainty of about 1 mm. An increase of the distance between the container and coil 1 by only 1.4 mm would explain the observed deviation of 50% between the calculated and the experimental values.
[28] The nonlinear Zeeman effect splits the resonance into $2F = 8$ resonances each separated by $\delta_{\text{NLZ}} = 2\nu_T^2/\nu_{\text{HFS}} = 851$ Hz where $\nu_{\text{HFS}} = 9.192631770$ GHz is the cesium hyperfine splitting.