ABSTRACT: To optimize sensitivity, there has been an increasing interest in the miniaturization of NMR detectors. In our lab, a stripline NMR detector has been developed, which provides high resolution and is scalable to a large range of sample volumes. These features make it an ideal detector for hyphenated techniques. In this manuscript, we demonstrate a stripline probe, which is designed for combining supercritical fluid chromatography (SFC) experiments with NMR. It features a novel stripline chip, designed to reduce the signal from the contact pads, which results in an improved lineshape. An external lock circuit provides stability over time to perform signal averaging or multidimensional experiments. As proof of concept, we demonstrate the SFC-NMR technique with this stripline probe using a mixture of cholesterol and cholestanol, which is relevant for studying cerebrotendinous xanthomatosis. Additionally, this probe makes it possible to record high-resolution spectra of samples with a high spin density. This means that it is possible to directly observe shifts due to the nuclear demagnetizing field in the "homomolecular" case, which is challenging using conventional probes due to broadening effects from radiation damping.

INTRODUCTION

In recent years, there has been an increasing interest in miniaturizing detectors for nuclear magnetic resonance (NMR) spectroscopy. This is because smaller coils can provide a higher sensitivity per spin for mass-limited samples. For example, for a flat circular coil, the sensitivity per sample volume is inversely proportional to its diameter, assuming the resistance of the coil does not change. Decreasing the coil diameter by a factor two doubles the signal-to-noise ratio of the NMR spectrum. Although the sensitivity increases, the number of spins that will fit inside the detected volume decreases. The size of the detector should therefore be matched to the available sample volume for optimal sensitivity. For mass- or volume-limited samples, such as body fluids, it is worthwhile to use a miniaturized detector. The sensitivity of the probe not only depends on its size but also on the coil design. Most of the research has been focused on developing planar helical coils and tightly wound solenoid microcoils. An alternative design is the planar microslot waveguide probe. However, with these types of microcoils, the resolution is often significantly reduced due to the close proximity of the microcoil structure to the sample. In our lab, a stripline detector has been developed, which solves this issue by using a planar structure oriented along the main magnetic field. The stripline NMR detector can record spectra with high sensitivity of samples contained within fused-silica capillaries. Also, the stripline can easily be designed for a wide range of sample volumes. This makes it straightforward to hyphenate this type of NMR detector to other techniques such as chromatography and capillary electrophoresis. An interesting development is the hyphenation of supercritical fluid chromatography (SFC) with stripline NMR.

In this manuscript, we demonstrate a novel NMR probe design, which is optimized for on-line SFC-NMR experiments. It features an improved stripline design, which provides a further increase in resolution, and an external lock circuit to provide stability over time for longer experiments. An interesting application of SFC-NMR is the separation and identification of lipids. In the work by Oostendorp et al., the analysis and quantification of lipids in body fluids are performed using 1H NMR without separation in a regular NMR setup. However, when too many lipids are present or when some are present at high concentration, identifying and...
quantifying the lipids can be difficult in these crowded NMR spectra. SFC-\(^1\)H NMR could resolve this problem since the lipids are separated before NMR analysis, thus solving the issue of a spectral overlap. We demonstrate the use of this novel probe design in a SFC-NMR setup for the separation and structural analysis of a mixture of cholesterol and cholestanol. This can be relevant for analyzing the blood plasma of patients suffering from cerebrotendinous xanthomatosis (CTX), which shows an accumulation of cholestanol due to a defect in a gene coding for the enzyme sterol 27-hydroxylase.

**EXPERIMENTAL SECTION**

**Split-Contact Stripline.** Although the stripline detector has a good homogeneity compared to other microcoil designs, the resonances often exhibited a broadening at the base. This is caused by the signal originating from the sample located on the contact pads (the top and bottom parts of the chip shown in Figure 1a). The magnetic field inhomogeneity in these areas is significant and can influence the sample's NMR behavior.

Figure 1. (a) Pictures of the regular stripline and (b) the new split-contact design. Both designs use ground planes (not shown) to confine the RF field. More information on the stripline design is provided in the Supplementary Information.

![Split-Contact Stripline](https://example.com/split-contacts.png)

Figure 2. Simulation of the distribution in \(B_1\)-field strengths in the sample as a function of position along the stripline for a regular stripline (blue contours) and a split-contact stripline (red contours). The black line shows the full width of the strip.

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![Simulated Field Distribution](https://example.com/simulated-field.png)

The expected difference in \(B_1\)-field strengths between the regular and split-contact stripline was simulated (COMSOL Multiphysics). The details of this simulation are provided in the Supplementary Information.

The magnetic field inhomogeneity in these areas is significant and can influence the sample’s NMR behavior. The expected difference in \(B_1\)-field strengths between the regular and split-contact stripline was simulated (COMSOL Multiphysics). The details of this simulation are provided in the Supplementary Information.
in this case, 15 μL of CDCl₃ (Cambridge Isotope Laboratories, 99.8 deuterated +0.05 v/v TMS), is introduced into the water flow to elute the sample. This plug flows toward the middle of the NMR stripline detector at which point the flow is stopped. Stopping the flow enables the acquisition of multiple scans to improve the signal-to-noise ratio. After the experiment, the valve returns to its original position and the sample flows back from the NMR probe to the waste. More details on the chromatography used in the SFC-NMR experiments are provided in the Supplementary Information.

For the NMR experiments, a home-built probe with the split-contact stripline as described above was used. The stripline chip used for SFC-NMR is of a larger design (dimensions are provided in the Supplementary Information) to accommodate a capillary with an outer diameter of 700 μm and an inner diameter of 500 μm, which means it has an active detection volume of ~1.6 μL. The SFC-NMR spectra and reference cholesterol spectrum were acquired with a relaxation delay of 10 s and an acquisition time of 1 s. For the SFC-NMR spectra, 5250 scans were recorded (16 h and 2 min) and 4500 scans for the reference cholesterol spectrum (13 h and 48 min, 25.9 mM). All spectra were recorded with a receiver bandwidth of 8 kHz. The spectra are referenced to TMS. The spectra were recorded in blocks of five scans. Before the summation of the spectra, the maxima of the TMS peaks was aligned to compensate for any remaining drift in the spectra over time, and the spectra were baseline corrected. All spectra were processed using ssNake.12

### RESULTS

**Split-Contact Stripline.** The split-contact stripline design was tested by comparing the RF-field strengths with a regular stripline design. For both designs, a chip was fabricated with a constriction length of 6.0 mm and a width of 500 μm. Using a capillary (OD 350 μm, ID 250 μm) with a 1 mm plug of water in FC-40, the power levels where calibrated such that the constricted part of the stripline chip produced an RF field of approximately 250 kHz. Afterward, the water sample was repositioned to the contact pad, and a nutation experiment was performed to determine the B₁-field produced at the contact pad. The positions of the sample are shown in Figure 3. The repositioning of the capillary did not influence the tuning and matching of the probe. The results of these experiments are shown in Figure 4. These B₁-fields correspond well to those predicted by the simulations. The regular stripline produced a B₁-field of approximately 21.5 kHz (simulated: 23.0 kHz) at the contact pad, whereas the split-contact stripline produced a B₁-field of only 6.0 kHz (simulated: 6.1 kHz). As can be seen from the figures, the sensitivity at the contact pads is also significantly lower for the split-contact design, as is expected from the reciprocity principle.13 The reduced B₁-field strength also means that when a pulse with a flip angle of 90° is given at the center of the stripline; the spins on the split-contacts experience a pulse with a flip-angle of 1.7° (compared to a flip-angle of 7.8° for a conventional stripline).

The reduced sensitivity of the contacts of the stripline results in an improvement of the resolution. This can be seen in Figure 5 where the CH₃ resonance of toluene is compared between a spectrum obtained using regular and split-contact striplines. The FWHM improves from 5.5 Hz to about 3 Hz; however, the biggest improvement can be seen in the base of the peak where most of the signal originates from the contact pads. The split-contact spectrum is still slightly asymmetric, which is most likely caused by local field disturbances, which are hard to correct using a regular shim system. An example of a source of these local field disturbances is the magnetic impurities in the “seed layer” used in the manufacturing of the stripline chip. One way to compensate for these local field disturbances would be to implement a Shim-on-Chip system.14

**External Lock.** To demonstrate the stability of the external lock with the stripline detector, we have recorded a series of single-pulse NMR spectra over time. In Figure S10, the drift of the CH₃ resonance of methanol is shown for a period up to 35 h. What can be observed is that without a lock, the magnetic field takes nearly 10 h to stabilize completely after the probe has been inserted into the magnet. Furthermore, there are sharp features visible at around 5, 15, and 30 h after the experiment was started. These are caused by the usage of the Bitter magnets in the High Field Magnet Laboratory (HFML) building about 100 m away from the NMR lab. To correct for these effects and for the drifting of the magnet, an external lock was implemented in the stripline probe. In this case, the external lock circuit contains a sample of deuterated acetone. Spectra of a sample of acetone were recorded for nearly 30 h. With the external lock, the initial drift after inserting the probe is corrected, as well as the influences of the magnets of the HFML. Some small variations in the resonance frequency can still be seen, with a maximum deviation of about 1 Hz. These variations are most likely caused by slight differences between the sample and lock sample.

**Demagnetizing Field.** An interesting effect that can be directly observed with this stripline probe is the shift of a resonance due to the nuclear demagnetizing field (long-range dipole–dipole interactions). The nonlinear effects of the nuclear demagnetizing field have been known for a long time, in particular, in the context of paramagnetic impurities,15,16 and the multiple echoes caused by the demagnetizing field have been observed experimentally in different materials.17

In this experiment, we placed a capillary with an inner diameter of 50 μm filled with H₂O in the stripline probe. The narrow diameter of the capillary minimizes radiation damping and lowers the B₁-gradient over the sample. Simple one-pulse NMR experiments were performed on this sample with varying pulse widths. The results are presented in Figure 6, which shows that there is a difference in the resonance frequency after a small flip angle compared to a near 180° pulse. This difference in frequency is about 2.3 Hz (3.8 ppb).
Due to the small diameter of the sample (i.e., 50 μm), the $B_1$ homogeneity is improved, which means that the resonance shift can be monitored for a number of cycles, as demonstrated in Figure 7a. In this figure, the black line approximately follows the resonance frequency. This line has an oscillation frequency (87 kHz) and a decay rate (Gaussian) identical to that of the signal amplitude (the orange line of Figure 7b) and only has a 90° phase difference. The resonance frequency also slowly shifts to lower frequency during the experiment, this is most likely caused by a slow increase in the sample temperature.

The effect of this demagnetizing field can be observed in this stripline setup because the sample is locked using an external lock circuit. In regular liquid-state NMR, it is common to lock on the deuterated solvent in the sample. This means that the demagnetizing field shift is not directly observed as shown here since the lock signal will be affected by the demagnetizing field also. It can also result in anomalous crosspeaks, for example, in COSY experiments.18

It is interesting to note that the magnitude (in Hz) of the shift in Figure 6 cannot be explained by the classical picture of bulk magnetization created by the sum of the individual proton spins. The shift between 0° and 180° in the classical picture is predicted to be 1.6 Hz,15,18 whereas we observe 2.3 Hz. In the quantum mechanical framework, the shift is described by long-range dipole–dipole interactions, which are not averaged by molecular motion.19 In this case, the predicted shift is $\frac{3}{2}$ times larger for “like” spins or so-called “homomolecular” spins.20 This is because the B-term in the dipolar Hamiltonian contributes to the total shift for these types of spins.

Direct detection of this kind of effect can only be done conveniently in a microliter setting. Recording spectra of samples with high spin densities in regular liquid NMR probes is usually complicated by other nonlinear effects, such as radiation damping. With the stripline probe, it is straightforward to reduce these effects so that the pulse width effect on the resonance position can be observed with high accuracy.

**SFC-NMR.** The stripline described above was optimized to analyze microliter (μL) sample volumes, thus matching the volumes coming from chromatographic techniques such as SFC. Therefore, we implemented the detector in our hyphenated SFC-NMR system. The design specifications of the stripline chip are provided in the Supporting Information. Due to high solubility of lipids in supercritical CO2, which has similar solvent strength as hexane, SFC is used more often in

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Figure 4. (a) Simulation and (b) experiment of the nutation frequency of a 1 mm plug of water in FC-40. The plug is positioned at the contact pads of a regular stripline (blue) and a split-contact stripline (orange).

Figure 5. CH$_3$ resonance of toluene recorded using a regular (blue) and a split-contact stripline.

Figure 6. Spectra of H$_2$O as a function of flip angle showing the shifting resonance. The black ellipse shows the path that the peak maximum/minimum makes. The width of the ellipse is 2.3 Hz.

Due to the small diameter of the sample (i.e., 50 μm), the $B_1$ homogeneity is improved, which means that the resonance...
the separation of different classes of lipids in the field of lipidomics. Here, we look at the SFC separation of cholesterol, one of the most known lipids, and cholestanol, which has a similar structure to cholesterol, differing in only one double bond (Figure 8). For the separation of these compounds, a 1 μL mixture of 28 mM cholestanol and 43 mM cholesterol in chloroform was injected into the SFC. The optimal separation conditions were obtained with a Torus Diol column from Waters, using a gradient of 2–5% methanol in CO₂ over 3 min, the result of which can be found in the Supporting information. Although the two molecules are very similar in structure, they can still be separated well under these conditions by SFC.

To test if the hyphenation of SFC-NMR could be a feasible analytic technique in the field of lipidomics, the split-contact stripline detector, optimized for μL volumes, was coupled in-line to the SFC apparatus. After SFC separation, both lipids were each collected six times in separate runs, before eluting them to the middle of the stripline NMR detector, in a plug of 15 μL of CDCl₃. There, the flow was stopped, allowing for the acquisition of multiple scans, to obtain a higher signal-to-noise ratio. To obtain a reference spectrum of cholesterol, instead of selecting the sample from the SFC, a 15 μL plug of 26 mM cholesterol in CDCl₃ was injected directly into the flow of water toward the NMR spectrometer. The resulting spectra are shown in Figure 8. By comparing the SFC-¹H NMR spectra with reference spectra, it is clear that the first chromatographic peak corresponds to cholestanol and the second peak corresponds to cholesterol. By using the split-contact stripline detector, the resolution and peak shape is improved (compared to using a regular stripline) and peak splittings due to J-couplings can be observed, e.g., around 3.55 ppm. The peak at 5.3 ppm, corresponding to the proton on the ring in cholesterol, which is absent in the cholestanol spectrum, confirms the selection of the correct molecules. Also, some shifts are observed for other peaks, reflecting the differences in the molecular structure. A mixture of very similar molecules can thus be separated and characterized by SFC-NMR. If the spectra are examined closely, a small impurity of cholesterol can however be observed in the cholestanol spectrum, probably due to a slight overlap of the peaks during chromatography. This does however not influence the
The SFC-NMR spectra shown here were acquired overnight over 16 h, which is a long time for an analysis. The signal-to-noise ratio is, however, much higher than required for the identification of these compounds. A spectrum with a signal-to-noise ratio of 3 for the peak around 5.3 ppm would require an experimental time of only 1 min. This is also due to the advantage of using the stripline probe with a larger volume. In our previous work, a stripline probe with a detection volume of only 150 μL was used. The current stripline has a volume of ~1.6 μL. The volume that is selected from the SFC is 100 μL for each selected peak. To get a higher sample concentration in the NMR detector, however, the mobile phase is evaporated out and the sample is redissolved in 15 μL of solvent, in this case, deuterated chloroform. The detection volume of 1.6 μL therefore matches the sample volume much better than in our previous probe in which only a small part of the sample was used for detection. The current probe volume could be increased even more, to better match the sample volume from the SFC. However, some mixing of the sample plug with the transportation medium, water, occurs at the edges of the plug. To prevent the water from interfering with the spectra, it is best not to use the full sample plug for detection but only the middle part. This μL volume stripline probe is therefore the best match in this hyphenation of SFC with in-line NMR detection. Although the absolute sensitivity of the stripline decreases by scaling up the design, the increased number of spins in the active detection volume leads to an overall gain in signal intensity. As an estimation, to obtain a signal-to-noise ratio of 3 for the cholesterol spectrum with the smaller volume stripline probe, 15 min of the experimental time is required, instead of 1 min. It is therefore beneficial to match the stripline probe volume to the sample volume obtained from the SFC.

The concentrations of cholestanol and cholesterol used in this SFC-NMR analysis are 28 and 43 mM, respectively, which are high compared to the concentrations of 0.2 and 3 mM that were used for the analysis technique for full structural characterization of lipids in body fluids. This type of stripline probe is well suited for hyphenated chromatography-NMR systems. The stripline described above was optimized to analyze μL sample volumes, thus matching the volumes coming from chromatographic techniques such as SFC. We therefore used the optimized stripline detector in our hyphenated SFC-NMR system.

An SFC-NMR setup has been demonstrated to separate a mixture of cholesterol and cholestanol with subsequent identification by NMR. These lipids are present in high concentrations in blood plasma of patients suffering from cerebrotendinous xanthomatosis (CTX). Both compounds can be isolated using this setup and analyzed with good resolution. In this proof-of-principle experiment, concentrations of cholesterol and cholestanol of 43 and 28 mM, respectively, were used for the analysis. However, with simple adaptations, like selecting the sample more often, this setup allows for the detection and identification of lipids in sub mM range, thus approaching the concentrations of 3 mM cholesterol and 0.2 mM cholestanol that are present in patients suffering from CTX. Although experiments still need to be performed on real patient samples, this is a proof-of-principle that the SFC-NMR setup with the optimized stripline probe can be a feasible analysis technique for full structural characterization of lipids in body fluids.
Complete contact information is available at:
https://pubs.acs.org/10.1021/acs.analchem.0c01827

Notes
The authors declare no competing financial interest.

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