Fluorine ($^{19}$F) MRI for Assessing Inflammatory Cells in the Kidney: Experimental Protocol

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Abstract

Inflammation is one underlying contributing factor in the pathology of acute and chronic kidney disorders. Phagocytes such as monocytes, neutrophils and dendritic cells are considered to play a deleterious role in the progression of kidney disease but may also contribute to organ homeostasis. The kidney is a target of life-threatening autoimmune disorders such as the antineutrophil cytoplasmic antibody (ANCA)-associated vasculitides (AAV). Neutrophils and monocytes express ANCA antigens and play an important role in the pathogenesis of AAV. Noninvasive in vivo methods that can quantify the distribution of inflammatory cells in the kidney as well as other organs in vivo would be vital to identify the causality and significance of inflammation during disease progression. Here we describe an noninvasive technique to study renal inflammation in rodents in vivo using fluorine ($^{19}$F) MRI. In this protocol we chose a murine ANCA-AAV model of renal inflammation and made use of nanoparticles prepared from perfluoro-5-crown-15-ether (PFCE) for renal $^{19}$F MRI.

This chapter is based upon work from the COST Action PARENCHIMA, a community-driven network funded by the European Cooperation in Science and Technology (COST) program of the European Union, which aims to improve the reproducibility and standardization of renal MRI biomarkers. This experimental protocol chapter is complemented by two separate chapters describing the basic concept and data analysis.

Key words Magnetic resonance imaging (MRI), Fluorine ($^{19}$F), Nanoparticles (NPs), Inflammation, Kidney, Mice

1 Introduction

Inflammation is one underlying contributing factor in the pathology of acute and chronic kidney disorders [1]. Studies also suggest that systemic inflammation can cause ischemic injury in a vital organ, such as the kidney, which could then result in repercussions in another distant organ downstream of the ischemic event, such as the heart [2–4]. Early inflammatory events governed by cells of the innate immune system, such as macrophages, probably promote
renal tissue injury but may also support repair [5–8]. Phagocytes such as dendritic cells (DC) and macrophages are considered to play a deleterious role in the inflammatory outcome of chronic kidney disease but may also contribute to organ homeostasis [9].

The kidney is often a target of systemic autoimmune disorders that are compounded by complex inflammatory processes [10]. Examples of life-threatening autoimmune disorders that affect the kidneys are the antineutrophil cytoplasmic antibody (ANCA)-associated vasculitides (AAV) manifesting as rapidly progressive necrotizing crescentic glomerulonephritis (NCGN) [11]. Neutrophils and monocytes express ANCA antigens and ANCA induces neutrophil extracellular traps that cause RIPK1-dependent endothelial cell (EC) damage via activation of the alternative complement pathway [12]. While the central role of neutrophil activation in ANCA-associated vasculitis and NCGN is clear, the role of monocytes/macrophages was only recently uncovered in a renal ANCA-associated vasculitis model [13].

Noninvasive in vivo methods that can quantify the level of inflammation in the kidney as well as other organs in systemic autoimmune disorders such as ANCA-associated vasculitides would be vital to identify the causality and significance of inflammation during the course of disease. One method to visualize inflammation by MRI makes use of MR contrast agents that modulate T2* and that are easily taken up by phagocytic inflammatory cells [14, 15]. Iron oxide particles including ultrasmall iron oxide agents (USPIO) have been used as susceptibility (T2*) MR contrast agents to target inflammatory cell populations. These particles are engulfed by phagocytic cells in the blood. Drawbacks of USPIO-based T2* studies include MR signal quantification and difficulty to distinguish contrast created by labeled cells from other intrinsic tissue contrasts [14].

Here we describe an alternative noninvasive technique to study inflammation in rodents in vivo using fluorine (19F) MRI. 19F MRI is performed in association with intravenous injections of perfluorocarbon (PFC) nanoparticles (NPs). These NPs are taken up by cells of the immune system traveling through the circulation into the inflammatory regions. Thus 19F MRI is ideal for studying distribution of inflammatory cell in vivo. In this protocol we chose a murine AAV model of renal inflammation that has been described in greater depth elsewhere [16] and made use of nanoparticles prepared from perfluoro-5-crown-15-ether (PFCE).

This experimental protocol chapter is complemented by two separate chapters describing the basic concept (please see the chapter by Waiczies S et al. “Functional Imaging Using Fluorine (19F) MR Methods: Basic Concepts”) and data analysis (please see the chapter by Starke L et al. “Data Preparation Protocol for Low Signal-to-Noise Ratio Fluorine-19 MRI”), which are both part of this book.

2 Materials

2.1 Animals

This experimental protocol is tailored for mice with a body mass of 20-30 g (e.g., wild type C57BL/6 mice or a disease model of renal inflammation). Here we describe briefly how to generate the AAV animal model. More thorough detail on the immunization, bone marrow transplantation as well purification of mouse MPO is given in the study establishing the MPO-AAV animal model [16]. Wild-type (WT) C57BL/6J mice (B6) (Jackson Laboratories, Bar Harbor, ME) and myeloperoxidase-deficient (MPO−/−) mice (generated by Aratani et al. [17]) were used in this protocol. MPO−/− mice were immunized with murine MPO at the age of 8–10 weeks, subjected to lethal irradiation, and then transplanted with MPO-expressing bone marrow cells. Animal experiments should be approved by animal welfare authorities and guidelines to minimize discomfort to animals (86/609/EEC).

2.1.1 Lab Equipment

1. NP preparation: Ultrasonic device with ultrasonic power (400 W) and frequency (24 kHz) for stand-mounted operation such as UP400S (Hielscher, Teltow, Germany) to prepare PFCE nanoparticles.

2. NP preparation: Titanium sonotrode for emulsifying samples from 5 to 200 ml (e.g., H3 from Hielscher, Teltow, Germany).

3. NP characterization: Dynamic light scattering instrument such as Zetasizer Nano (Malvern Instruments, Malvern, Worcestershire, UK) to characterize particle size (Z-average) and polydispersity index (PDI) of the PFCE nanoparticles.

4. NP application: Mouse restrainer for intravenous administering the PFCE nanoparticles.

5. Anesthesia: Isoflurane inhalation system which can adjust different levels of isoflurane such as Isoflurane Vapor 19.1 (Draeger, Draegerwerk, Luebeck, Germany). The range of isoflurane level that is used for anesthesia in mice is 0.5–1.5%. Please refer to the chapter by Kaucsar T et al. “Preparation and Monitoring of Small Animals in Renal MRI” for an in-depth description and discussion of the anesthesia.

6. Anesthesia: Mouse chamber connected to isoflurane inhalation system and gas-mixing system to anesthetize mice by inhalation narcosis prior to transfer to the MR scanner.

7. Gases: O2, and compressed air, as well as a gas-mixing system such as FMI (Förhr Medical Instruments GmbH, Seeheim-Ober Beerbach, Germany) to achieve the required physiological O2/air mixture in combination with the isoflurane gas.
2.2 MRI Hardware

The general hardware requirements for renal $^1$H MRI on mice and rats are described in the chapter by Ramos Delgado P et al. “Hardware Considerations for Preclinical Magnetic Resonance of the Kidney.” The technique described in this chapter was tailored for a 9.4 T MR system (Biospec 94/20, Bruker Biospin, Ettlingen, Germany) but advice for adaptation to other field strengths is given where necessary.

1. $^1$H/$^{19}$F dual-tunable volume RF coil (35 mm inner diameter, 50 mm length; Rapid Biomed, Würzburg, Germany).
2. $^1$H/$^{19}$F dual-tunable volume RF coil (18 mm inner diameter, 39 mm length) for ex vivo imaging [18].
3. $^{19}$F cryogenic quadrature RF surface probe ($^{19}$F-CRP) operated at ~28 K (see Note 1).
4. A physiological monitoring system that can track respiration and temperature during the MR procedure such as the Monitoring & Gating System and PC-sam software from SA Instruments (SAII, Stony Brook, NY, USA).
5. Mouse sled with a breathing mask connected to the isoflurane system.

2.3 MRI Techniques

Typically, $^{19}$F MR studies applying PFC compounds such as PFCE to study inflammation in vivo employ the turbo spin echo (SE) rapid acquisition using relaxation enhancement (RARE) sequence [18–22]. This method reduces acquisition time by accumulating multiple echoes within a single repetition time [23]. Typically $T_1$ of these $^{19}$F compounds is in the range of 0.5–3 s, depending on the compound and also magnetic field strength ($B_0$). However, if employing paramagnetic macrocyclic PFC compounds complexed to lanthanides, $T_1$ values can be reduced to the order of 1–15 ms and $T_2^*$ values correspondingly reduced to 0.4–12 ms and a radial zero echo time (ZTE) sequence might be better suited [24].

1. $^1$H MR sequence: 2D and 3D FLASH protocols are standard sequences on Bruker MRI systems, where they are called “FLASH_2D”, “FLASH_3D” or “1_Localizer_multi_slice” (see Note 2).
2. $^{19}$F spectroscopy sequence: Block pulse for nonlocalized (global) is a standard sequence on Bruker MRI systems, where it is called “SINGLEPULSE” (see Note 3).
3. $^{19}$F MR sequence: 3D RARE protocol for measurements. This is a standard sequence on Bruker MRI systems, where it is called “TurboRARE_3D” in Paravision 5, “T2_TurboRARE_3D” in Paravision 6 (see Note 4).
Typically $T_1$ of PFC compounds is in the range of 0.5–3 s, depending on the compound and also magnetic field strength ($B_0$). When working with a standard diamagnetic PFC such as PFCE that is documented in the literature, it is recommended that the $T_1$ and $T_2$ of the compound be studied at 37 °C before starting with the first in vivo experiments to study inflammation. According to the measured relaxation times ($T_1/T_2$), the optimal settings for RARE, namely echo train lengths (ETL) and repetition time (TR) should be calculated to improve sensitivity thresholds.

1. **Repetition time (TR):** for RARE, TR will be limited by ETL and the number of slices. A short TR is desirable for SNR efficiency (0.8–1.5 s depending on $T_1$) keeping in mind that $T_1$ will change with changes in oxygenation status (see Note 5).

2. **Flip angle (FA):** for the block pulse and RARE sequence, the FA for the excitation pulse should be at 90°. Additionally the RARE sequence has a refocusing pulse with a FA of 180°. To maximize the SNR and contrast, the flip angle has to be separately calculated for FLASH considering the Ernst angle:
   \[ \alpha_E = \cos^{-1}\left(e^{-\Delta T_1/T_1}\right) \]

3. **Echo train length (ETL):** in RARE, use a value as high as possible to reduce scan time. In the TurboRARE sequence on Bruker MRI systems, ETL is referred to as “rare factor”.

4. **Echo time (TE):** use the shortest effective TE and echo spacing ($\Delta T_E$) possible. Especially for very high ETL, TE can become very long. Centric encoding can be used to reduce the effective TE, generally at the cost of introducing minor image artifacts.

5. **Acquisition bandwidth (BW):** long enough to shorten the $\Delta T_E$ without compromising the SNR, which decreases with increasing BW as a result of increased noise level.

6. **Geometry:** Adapt so that whole abdomen fits into FOV in L-R direction (approx. 25 mm for mice) and use frequency encoding in H-F- direction. Choose a 3D volume suited to cover the whole body width and as much of the length of the mouse that the RF coil can afford as possible. With a typical mouse body resonator as the one used in this protocol one could use a FOV of $60 \times 30 \times 30$ mm and matrix size of $400 \times 200 \times 200$ (for $^1$H) and $128 \times 64 \times 64$ (for $^{19}$F).

7. **Resolution/acceleration:** Use the highest in-plane resolution that the SNR allows, typically around 100–150 μm for $^1$H MR scans and 250–500 μm for $^{19}$F MR scans. Zero-filling in phase encoding direction can be helpful to speed up acquisition while increasing the number of averages to improve SNR.
especially for $^{19}$F MR scans. One may use half Fourier in read direction (asymmetric echo) to further shorten the first TE. Reducing the excitation pulse length to below 1 ms would then also help to shorten TE.

8. For an example of parameters used in this chapter, please see Notes 7 and 13.

3.2 $^{19}$F Nanoparticle Preparation, Characterization, and Application

1. **NP preparation**: Emulsify 1.2 M PFCE (Fluorochem, Derbyshire, UK) in Pluronic F-68 (Sigma-Aldrich, Germany) for 10 min using a cell disrupting titanium sonotrode connected to an ultrasonic device and employing a continuous pulse program for 60 s. Use ear protection while sonicating the mixture to avoid hearing damage and loss.

2. **NP characterization**: Measure mean particle size (in nm), polydispersity index (PdI), and zetapotential (mV) by using a dynamic light scattering machine such as the one listed above. Use the $z$-average diameter for particle size since it gives an intensity-weighted harmonic diameter and is ideal for comparing different analyses. The nanoparticles should have a PdI < 0.3 indicating a relatively low polydispersity and narrow size distribution (see Note 8).

3. **NP application**: Administer $^{19}$F nanoparticles via tail vein at a dose of 5–80 µmol of PFCE molecules, depending on the frequency of the bolus injections. Start administering $^{19}$F nanoparticles at relevant time-points of your inflammatory model, for example, in MPO immunized MPO$^{-/-}$ mice subjected to lethal irradiation we started intravenous application of PFCE nanoparticles 4–8 weeks following transplantation of MPO-expressing bone marrow cells (see Note 9).

3.3 Preparation Prior to $^{19}$F/$^1$H MRI Scans

Four to 18 h following the last intravenous administration of $^{19}$F nanoparticles prepare the mice for $^{19}$F/$^1$H MRI:

1. First anesthetize the mice by inhalation narcosis using a mouse chamber connected to a isoflurane inhalation system and gas-mixing system (see Note 10).

2. Adjust the flow rate for air and O$_2$ at 0.2 and 0.1 l/min respectively and 3% isoflurane (adjusted from a vaporizer) for about 2 min until the required level of anesthesia is reached (no response following toe pinch).

3. Transfer mice to the MR scanner. Should a quantification of inflammation be required, a reference tube with a known concentration of $^{19}$F nanoparticles should be placed in proximity to the region of interest. For quantification of signal please refer to the chapter by Starke L et al. “Data Preparation Protocol for Low Signal-to-Noise Ratio Fluorine-19 MRI.”
4. While keeping the flow rate for air and O₂ constant, adjust the isoflurane vaporizer to 0.8–1.5% until an optimal breathing pattern is reached.

5. Set up the temperature monitoring (rectal probe) and respiratory monitoring (balloon on chest) unit. A respiratory rate of 70–90 breaths per minute is recommended. Keep the body temperature at 36–37 °C during the experiment by employing a warm water (or alternatively warm air) circulation system.

6. Tune the RF coil to both the ¹H resonance frequency (e.g., 400.1 MHz for 9.4 T) and to the ¹⁹F resonance frequency (e.g., 376.3 MHz for 9.4 T) and match the characteristic impedance of the coil to 50 Ω using the tuning monitor of the animal MR scanner.

7. Perform anatomical imaging as described in the chapter by Pohlmann A et al. “Essential Practical Steps for MRI of the Kidney in Experimental Research.” Set up a 2D FLASH protocol for the acquisition of anatomical kidney ¹H scans (see Note 7).

8. Perform localized shimming on the kidney imaging as described in the chapter by Pohlmann A et al. “Essential Practical Steps for MRI of the Kidney in Experimental Research” (see Note 11).

9. Save the parameters of all adjustments (such as iterative shimming calculations and reference power values) performed during the first ¹H scans for application into the ¹⁹F scans.

Following acquisition of the anatomical ¹H kidney scans, in vivo ¹⁹F MR images of the kidney can be acquired and later overlaid onto the ¹H MR images. An example of an in vivo ¹H/¹⁹F MRI is shown in Fig. 1.

1. Load the SINGLEPULSE FID-sequence with a TR of at least 1000 ms.

2. Set nucleus to ¹⁹F (e.g., in “Open Edit Scan” in Bruker’s Paravision 5.1 or “System” tab in Bruker’s Paravision 6). Since the ¹⁹F MR signal is too low for automatic adjustments, apply the same settings used for the ¹H anatomical scans (see Note 12).

3. Deselect the automatic reference gain (RG) and set on maximum (e.g., in Edit Method in Bruker’s Paravision 5.1 or using the “Instruction” tab (GOP) and “Setup” tab in Paravision 6).

4. Start the SINGLEPULSE sequence using setup mode. If the ¹⁹F spectral signal within the acquisition-window is too low, add more averages until a signal is clearly visible. Adjust the basic frequency in order to center the ¹⁹F spectral peak at 0 Hz in the acquisition window. Apply this basic frequency, press Stop and apply.
5. Setup a TurboRARE 3D protocol for the \textsuperscript{19}F scans.

6. Use the same geometry used for anatomical \textsuperscript{1}H imaging but reduce the matrix size for increased SNR and use a rare factor of at least 32. Set nucleus to \textsuperscript{19}F. Deselect automatic adjustments (as above). For an example of parameters, please see Note 7.

7. When the scans are finished retract the mouse-holder from the MR scanner. Disconnect the mouse carefully from the holder. If the mouse is not sacrificed for ex vivo analysis (e.g., high resolution \textsuperscript{19}F MRI of the kidney, see below) following the MR scans, closely monitor until it has completely recovered from anesthesia. Body temperature regulation might still be affected after the anesthesia, so during the recovery process, put the mouse in a separate cage that is placed on a warm temperature regulated pad. Once the mouse has completely recovered from anesthesia, you may return it to its holding cage and to the animal room.

Kidney inflammation can also be studied with high spatial resolution ex vivo \textsuperscript{19}F MRI, for example, by using a transceive \textsuperscript{19}F CryoProbe, which we previously used to study brain inflammation in a model of CNS autoimmunity [25]. An example of a high resolved ex vivo \textsuperscript{1}H/\textsuperscript{19}F MRI of the kidney is shown in Fig. 2.

1. At the end of the in vivo experiment, anesthetize mice with a terminal dose of ketamine and xylazine. Ensure the required level of anesthesia is reached (no response following toe pinch).

2. Transcardially perfuse mouse with 20 ml PBS followed by 20 ml 4% paraformaldehyde.
3. Harvest relevant organs (e.g., kidney, liver, and spleen).

4. Transfer organs to a container filled with 4% PFA and store at 4 °C.

5. Prior to the high resolution 19F MRI of the kidney, embed the kidney in 1% low melting agarose in a 1.5 ml Eppendorf tube.

6. 19F-CRP adjustments: since the 19F signal is too low for automatic adjustments, use a 19F calibration phantom (fill highly fluorinated substance such as trifluoroethanol with water in a 1.5 ml Eppendorf tube. Place the 19F calibration tube under the 19F-CRP and repeat steps 13–16 to calculate the working frequency. Load and run a 1_Localizer_multi_slice sequence (see above). Adjust the reference power using the adjustments platform: select a coronal slice of 2 mm thickness and place it close to the RF coil’s surface. Select an initial power two orders of magnitude smaller than what is usually expected and press start. Run the rest of the adjustments using the same sequence. Save the shim settings.

7. 19F-CRP imaging: place the ex vivo kidney embedded in a 1.5 ml Eppendorf tube under the 19F-CRP surface. Run the working frequency adjustments as per 13–16, load the shim
calculations, and adjust the reference power as described above. Set the RG to maximum. Load a 2D FLASH protocol and set it up for $^{19}$F imaging. Examples for high and medium resolution $^{19}$F imaging are given in Note 13.

8. Set up a 2D FLASH protocol for the acquisition of anatomical kidney $^{1}$H scans. Use the same geometry used for $^{19}$F imaging (see Note 13).

4 Notes

1. The transceive $^{19}$F cryogenic quadrature RF surface probe ($^{19}$F CryoProbe) operates at ~28 K with a dual cooled preamplifier at the base running at ~77 K. It has a similar geometry to the existing Bruker $^{1}$H quadrature CryoProbes. More details on the $^{19}$F CryoProbe are available in our previous study [25].

2. The FLASH_3D protocol is especially adapted for whole-body imaging of mice (the gradient-system and the volume-resonator need to have a linear region of about 8 cm).

3. The block pulse program (SINGLEPULSE) contains the necessary elements for a simple transmit/receive experiment, sending a pulse and acquiring an FID afterward.

4. A 2D version of this sequence is also available, which allows thicker slices for a general overview but suffers from low SNR for most in vivo applications.

5. Special attention should be given when studying inflammation in models where tissue oxygen levels are likely to change, for example, following ischemic events. In these cases, $T_1$ weighting needs to be reduced at cost of SNR efficiency by increasing $TR \gg T_1$ (typically $TR = 3–5 \times T_1$).

6. When using transmit-receive surface coils a $B_1$ correction should be considered in order to compensate for the intrinsic spatial gradient in coil sensitivity ($B_1^-$) and excitation field ($B_1^+$ inhomogeneity), which results in signification variation in the excitation FA, decreasing with increasing distance from the RF coil surface. This severely reduces image homogeneity and hampers the acquisition of the absolute signal intensity values for $^{19}$F quantification techniques.

7. Example for a 30 g mouse at 9.4 T and FOV of $50 \times 25 \times 25$ mm, (1) $^{1}$H 2D FLASH protocol: $TR = 579.4$ ms, $TE = 5$ ms, FA = 75, matrix = 256 $\times$ 128. (2) $^{19}$F 3D RARE protocol: $TR = 800$ ms, $TE = 6.16$ ms, Matrix = 256 $\times$ 128 $\times$ 128, RARE Factor = 32–64.
8. The polydispersity index (PdI) is extrapolated from the DLS function and quantitatively describes the particle size distribution best. PdI ranges from 0.01 for monodispersed particles to 0.7 for particles that have a very broad size distribution. The $z$-average diameter gives the mean diameter based on intensity of scattered light and sensitive to presence of large particles, peak diameter, peak width, and PdI.

9. In MPO-AAV mice we administered one bolus of PFCE nanoparticles intravenously (80 $\mu$mol in 100 $\mu$l) 8 weeks following transplantation of bone marrow cells.

10. Mice can be alternatively anesthetized with an intraperitoneal injection of ketamine and xylazine.

11. Shimming is particularly important, since macroscopic magnetic field inhomogeneities affect the exact resonance frequency of the PFCE compounds and might affect quantification of the $^{19}$F MR signal. Shimming should be performed on a voxel enclosing both kidneys using either the default iterative shimming method or the Mapshim technique (recommended). However, the Mapshim technique is not available for X-nuclei-only RF coils. An alternative in this case is to use a highly fluorinated sample to calculate the shims.

12. Since the NMR properties of $^{19}$F and $^1$H are similar, the same MR setup and MR parameter settings can be used for both nuclei.

13. Example for ex vivo high-resolved $^{19}$F MRI at 9.4 T using the $^{19}$F-CRP and a 2D FLASH: TR = 11 ms, TE = 2.7 ms, FA = 30, Avg = 2250, Repetitions = 35, FOV = 20 $\times$ 20, matrix = 256 $\times$ 256, 1 mm slice thickness. Example for ex vivo low-resolved $^{19}$F MRI at 9.4 T using the $^1$H/$^{19}$F dual-tunable volume RF coil (18 mm inner diameter) and a 3D RARE method: TR = 800 ms, TE = 5 ms, Avg = 256, Repetitions = 13, FOV = 60 $\times$ 30 $\times$ 30, matrix = 64 $\times$ 32 $\times$ 16. $^1$H 2D FLASH protocol: TR = 18.7 ms, TE = 5.5 ms, FA = 25, matrix = 171 $\times$ 256, 1 mm slice thickness.

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References


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