TECHNICAL NOTE

A robust broadband fat-suppressing phaser T₂-preparation module for cardiac magnetic resonance imaging at 3T

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Purpose: Designing a new T₂-preparation (T₂-Prep) module to simultaneously provide robust fat suppression and efficient T₂ preparation without requiring an additional fat-suppression module for T₂-weighted imaging at 3T.

Methods: The tip-down radiofrequency (RF) pulse of an adiabatic T₂-Prep module was replaced by a custom-designed RF-excitation pulse that induces a phase difference between water and fat, resulting in a simultaneous T₂ preparation of water signals and the suppression of fat signals at the end of the module (a phaser adiabatic T₂-Prep). Numerical simulations and in vitro and in vivo electrocardiogram (ECG)-triggered navigator-gated acquisitions of the human heart were performed. Blood, myocardium, and fat signal-to-noise ratios and right coronary artery vessel sharpness were compared against previously published adiabatic T₂-Prep approaches.

Results: Numerical simulations predicted an increased fat-suppression bandwidth and decreased sensitivity to transmit magnetic field inhomogeneities using the proposed approach while preserving the water T₂-Prep capabilities. This was confirmed by the tissue signals acquired in the phantom and the in vivo images, which show similar blood and myocardium signal-to-noise ratio, contrast-to-noise ratio, and significantly reduced fat signal-to-noise ratio compared with the other methods. As a result, the right coronary artery conspicuity was significantly increased.

Conclusion: A novel fat-suppressing T₂-Prep method was developed and implemented that showed robust fat suppression and increased vessel sharpness compared with conventional techniques while preserving its T₂-Prep capabilities.

KEYWORDS
3T magnetic resonance imaging (MRI), adiabatic, angiography, coronary, fat suppression, noncontrast, T₂ preparation

INTRODUCTION

Noncontrast-enhanced pulse sequences often lack the required contrast to distinguish blood from the myocardium in cardiac MRI. T₂-preparation (T₂-Prep) modules¹ allow this contrast to be increased by using the difference in T₂ relaxation times between these two tissue types.²,³ Such modules are also used for robust T₂ mapping of the myocardium and knee cartilage.⁴-⁶ In these applications, unwanted signal from fat can compromise the delineation and anatomical
visualization of the tissue of interest, such as cartilage, myocardium, or the coronary vessels, and thus decrease image quality. Unsuppressed fat signal may also lead to water-fat signal cancellation and can accentuate artifacts.

To suppress an unwanted lipid signal, pulse sequences usually include additional fat signal-suppressing approaches, such as chemically selective saturation (CHESS), water-excitation, or frequency-selective inversion RF pulses. Higher magnetic field strengths may complicate conventional fat-saturation and T2-Prep techniques, which are sensitive to B0 and B1 field inhomogeneities. The use of adiabatic RF-excitation pulses addressed the sensitivity to B1 inhomogeneities. By introducing a delay before the tip-up pulse of an adiabatic T2-Prep ion module, the saturation of spins was able to resonate at a target frequency. However, the fixed fat-suppression bandwidth may not be sufficient to cope with B0 inhomogeneities around the heart. Decreasing the bandwidth of the tip-down RF pulse attenuated off-resonance fat signals but depended on a precise RF-excitation angle, thus being sensitive to B1 inhomogeneities while still requiring an additional fat-saturation module. Moreover, the increase in RF-excitation angle of the tip-down and tip-up pulse, required to negate inversion recovery of the fat signal in-between the T2-Prep and the start of the acquisition, reduced the efficiency of the T2 preparation of on-resonance water.

The goal was to develop a novel T2-Prep approach with a dephasing tip-down RF pulse that addresses the aforementioned challenges of robustness to B0 and B1 inhomogeneities while preserving T2 contrast and simultaneously suppressing fat. The sensitivity of this technique to B0 and B1 inhomogeneities, as well as the effect of changing T2-Prep durations and T1 and T2 relaxation times, was quantified by numerical simulations. The simulation results were validated in phantom and electrocardiogram (ECG)-triggered navigator-gated acquisitions of the human heart in healthy volunteers and were compared against routinely used techniques.

2 METHODS

2.1 Tip-down RF pulse design

The tip-down RF excitation pulse of an adiabatic T2-Prep module was replaced by a custom-designed pulse (Figure 1A). The purpose of the RF design is to rotate both on- and off-resonance spins into the transverse plane while introducing a specific phase difference (Δφ) between the two spin populations at the T2-Prep echo time (Figure 1B). Then, the nonselective tip-up pulse rotates the on-resonance magnetization back along the longitudinal axis, but because of the phase difference Δφ, the off-resonance spins are rotated into the transverse plane and spoiled by subsequent spoiler gradients (Figure 1C). The proposed fat-suppressing phaser adiabatic T2-preparation (PA-T2-Prep) has a bandwidth that comprises both water and fat (Figure 1D), and Δφ can be adjusted to obtain the desired longitudinal components of the off-resonance magnetization at the end of the module, without altering the T2 preparation of on-resonance spins and without increasing the RF power. Fat magnetization is rotated with each applied RF pulse, which increases the robustness to B1 inhomogeneities.

The RF profile of the tip-down pulse was approximated as the inverse Fourier transform of two rectangular functions representing the desired transverse magnetization of water and fat. One function represents the magnetization with frequencies below f0 (fat) and bandwidth BW/2 that will be spoiled after T2 preparation. The other represents the magnetization with frequencies above f0 and bandwidth BW/2 that will be T2 prepared (Figure 1D). In addition, a phase of ±Δφ/2 is introduced to make the total phase difference Δφ (Figure 1D). Following an inverse Fourier transform, the pulse shape is proportional to (Figure 1E): 

$$\frac{1}{2} \sin \left( \frac{2\pi BW/2}{2\pi} \right) \left( e^{i\Delta \phi/2} + e^{-i\Delta \phi/2} \right)$$

where sinc is the normalized cardinal sine function and t is time. The pulse duration PD was set to 8180 µs, the upper limit imposed by the vendor-provided source code. The pulse bandwidth was chosen to have a fat-suppression bandwidth close to 700 Hz (total bandwidth 1400 Hz) while having zero-crossings at the beginning and the end of the pulse, which is satisfied if $$BW = \frac{2k}{PD}, k \in \mathbb{Z}^+.$$ BW was set to 1467 Hz (ie, k = 3), the RF-excitation angle was kept to 90°; the pulse off-resonance frequency ($f_0$) was set to −100 Hz, in-between water (0 Hz) and fat (−407 Hz). The optimal Δφ for water-fat–phase separation, facilitating fat suppression, was initially determined using simulations and subsequently fine-tuned in a phantom. To compensate for phase gain based on $f_0$ and Δφ, the initial phase of the phaser tip-down pulse was set to $-\frac{1}{2} (PD2\pi f_0 + \Delta \phi)$.

Although a small flip-angle approximation was used for the PA-T2-Prep design, a comparison with Bloch simulations shows negligible impact (<5°) on the amplitude and phase of the magnetization (Supporting Information Figure S1).

2.2 Numerical simulations

The effects of the conventional adiabatic T2-prep (CA-T2-Prep$^{16}$ with CHESS fat saturation (+FS) and the proposed PA-T2-Prep on the longitudinal magnetization were quantified using numerical simulations performed in MATLAB (The MathWorks). The tip-down and tip-up RF pulses were modeled as Hamming-windowed sinc functions of 800 µs of opposite polarity for the CA-T2-Prep+FS. For the
PA-T2-Prep, the tip-down pulse as described above was used, whereas the tip-up pulse was identical to that of the CA-T2-Prep. The two hyperbolic secant refocusing adiabatic pulses were modeled using their RF-pulse shapes as implemented in the sequence with an excitation bandwidth of ±800 Hz. The CHESS pulse was modeled as implemented in the sequence, a 5120 µs, −407 Hz Gaussian pulse with RF excitation angle = 100°. The simulations were performed by Euler integration of the Bloch equations using steps of 1 µs. The $T_1$ and $T_2$ relaxation times were set to those of fat at 200 ms and 50 ms, the off-resonance frequency was varied from −1000 Hz to 1000 Hz, and the $B_1$ amplitude ranged from 0% to 200% to account for $B_0$ and $B_1$ inhomogeneities. Magnetization was considered suppressed if its longitudinal component at the end of the T2-Prep was reduced to between ±10% of the starting magnetization $M_0$. The excitation bandwidth reflects

\[ M_{xy}(f) = \begin{cases} e^{-\frac{\Delta \phi}{2} i} & \text{if } -\frac{BW}{2} \leq f < 0 \\ e^{+\frac{\Delta \phi}{2} i} & \text{if } 0 \leq f < +\frac{BW}{2} \\ 0 & \text{otherwise} \end{cases} \]

\[ B_1(t) \propto \text{sinc}\left(\frac{BW}{2} t\right) \cos\left(\frac{\pi}{2} B_1 A_{\text{IFT}} t + \Delta \phi / 2\right) \]

**Figure 1** Diagram of the proposed phaser T2-preparation module and design of the phaser tip-down radiofrequency (RF) pulse. A, Overview of RF pulses used in the T2 preparation module. The module is followed by a spoiling gradient (not illustrated). B, Excitation of the water (blue) and fat (yellow) magnetizations by the phaser tip-down pulse. The solid and dashed vectors represent the magnetization before and after the excitation, respectively, and the black-dashed trajectories represent their rotations. Both fat and water are rotated into the transverse plane, but the pulse introduces a phase difference $\Delta \phi$ between the two. C, Excitation of the water and fat magnetizations by broadband tip-up pulse. The T2-prepared–water magnetization is restored along the longitudinal axis whereas the fat is kept near the transverse plane because of $\Delta \phi$. D, Desired transverse magnetization fraction $|M_{xy}(f)|$ (red) and magnetization phase $\arg(M_{xy}(f))$ (green) after the phaser tip-down RF excitation as function of spin off-resonance frequency. BW: RF-excitation bandwidth, $\Delta \phi$: phase difference between fat and water magnetization, $f_0$: RF-excitation–pulse frequency. E, $B_1$ amplitude of the phaser tip-down RF-excitation pulse as a function of time, calculated as the inverse Fourier transform of (D). A Bloch simulation of the RF-pulse profile (1E), as well as the small flip-angle approximation, can be found in Supporting Information Figure S1.
the region of longitudinal magnetization being ±10% from the on-resonance magnetization at 100% B1 amplitude. Note that 100% M0 is not possible because of the T2-Prep module.

Additional simulations were performed to investigate the effect of different T2-Prep durations and Δφ on the longitudinal magnetization of blood (T1/T2 = 1463/221 ms), myocardium (T1/T2 = 1072/31 ms), and fat (T1/T2 = 211/42 ms) compartments with relaxation times determined in our "heart" phantom to enable a comparison with experimental data. A 6-peak fat model was used with frequency components of −469 Hz, −420 Hz, −321 Hz, −239 Hz, −48 Hz, and 74 Hz with respective amplitudes of 0.087, 0.694, 0.128, 0.004, 0.039, and 0.048.

2.3 Phantom studies

A three-compartment cylindrical heart phantom was used that contains mixed solutions of agar and NiCl2 (Sigma-Aldrich) or baby oil (Johnson & Johnson), mimicking the magnetic relaxation properties of blood (T1 = 1463 ± 24 ms, T2 = 221 ± 15 ms), muscle (T1 = 1072 ± 8 ms, T2 = 31 ± 0.4 ms), and fat (T1 = 211 ± 1 ms, T2 = 42 ± 0.3 ms). Acquisitions were performed on a clinical 3T scanner (MAGNETOM Prisma; Siemens Healthcare) using an 18-channel spine and 16-channel chest RF coil with a segmented three-dimensional Cartesian gradient echo sequence with a field of view of 160 × 96 × 95 mm³ and resolution of 1 × 1 × 5 mm³. After the T2-Prep with 10-ms hyperbolic secant adiabatic refocusing pulses, 25 k-space lines were acquired using centric ordering: pulse repetition time = 5 ms, echo time = 2.5 ms, RF excitation angle = 15°, bandwidth = 501 Hz/pixel, and time between T2-Preps = 1 s. The T2-Prep duration was varied from 40 ms to 60 ms to 80 ms and the Δφ was varied between 90° and 270° by steps of 5° (randomized acquisition order), and the resulting average signal from each of the three compartments was quantified. The Δφ yielding the lowest fat signal for a T2-Prep duration of 40 ms was used for the rest of the study.

Additional experiments were performed in a National Institute of Standards and Technology phantom to investigate the effect of a range of T1 and T2 values, T2-Prep durations, and Δφ values on the signal behavior in each compartment.

Mean compartment signals were quantified after using five different T2-Prep modules: no T2-Prep, CA-T2-Prep, CA-T2-Prep+FS, PA-T2-Prep, and the water-selective adiabatic T2-Prep with fat saturation (WSA-T2-Prep+FS). The RF-excitation angle of the CHESS pulse was set to 100°, which was chosen based on experimental fine-tuning performed in the heart phantom. The B₀ and B₁ fields were intentionally not shimmed to test the sensitivity of each technique to inhomogeneities. Scan–rescan acquisitions were performed for quantification of blood, myocardium, and fat-compartment mean signal and noise using the acquisition-subtraction method. A 1-h–long scan with 100 averages was acquired prior to the scan–rescan experiment to reach thermal steady state for true noise quantification. A compartment-specific SNR was computed relative to the no T2-Prep case to emphasize compartment-specific differences across T2-Preps. A B₀ map with a ±1000 Hz bandwidth was computed from two acquisitions with the same acquisition parameters as described previously, without T2-Prep and with an echo time of 2.5 ms and 3.0 ms. Scaling in B₀ maps was cropped to ±500 Hz. Additionally, a B₁ map from two acquisitions with RF-excitation angles of 60° and 120° was computed using a double-angle method and a pulse-repetition time of 10 s.

2.4 In vivo study

Human studies were approved by the local ethics committee. Whole-heart free-breathing ECG-triggered navigator-gated scans were acquired in healthy volunteers who gave written and informed consent (N = 6, age = 27 ± 4 years). To keep the scan time below 1 h per participant, only the CA-T2-Prep, CA-T2-Prep+FS, and PA-T2-Prep were tested. The same imaging sequence was used as described earlier, except for a field of view of 230 × 368 × 86.4 mm³ and an isotropic pixel size of 1.2 mm. Images were reformatted using Soap-Bubble, a semiautomated reformatting and vessel-tracking software package. The right coronary artery (RCA) vessel sharpness was computed in a 4-cm segment using Soap-Bubble. Blood, myocardium, chest fat, and epicardial SNR were quantified in chest fat, epicardial fat, blood, and myocardium. The SNR was approximated by dividing the average signal from regions of interests drawn in respective tissues by the standard deviation of the background noise. The CNR was quantified between blood and myocardium compartments, as well as blood and epicardial fat and myocardium and epicardial fat. Differences between the CA-T2-Prep and PA-T2-Prep, as well as between the CA-T2-Prep+FS and PA-T2-Prep, were tested via a Student’s t test for paired data. A Bonferroni correction was applied to correct for multiple comparisons: P values obtained with individual t tests were multiplied with the amount of comparisons to keep the significance threshold at P < .05. Additionally, the specific absorption rate (SAR) was recorded from the system console.

3 RESULTS

3.1 Numerical simulation

The PA-T2-Prep simulation (Figure 2A) predicted an increased robustness in fat suppression against B₀ and B₁ field inhomogeneities in comparison with the CA-T2-Prep+FS...
For an off-resonance frequency of $-407$ Hz, the magnetization was suppressed if $B_1$ was between 84% and 111% for the CA-T2-Prep+FS versus 75% and 130% for the PA-T2-Prep. At a $B_1$ amplitude of 100%, the fat-suppression bandwidth was 162 Hz for the CA-T2-Prep+FS and 627 Hz for the PA-T2-Prep (Figure 2C,D). At 100% $B_1$ amplitude the on-resonance magnetization was approximately 54% and approximately 51% for the CA-T2-Prep+FS and PA-T2-Prep, respectively (Figure 2C,D). On-resonance magnetization was excited if $B_1$ was between 60% and 125% the CA-T2-Prep+FS versus 67% and 127% for the PA-T2-Prep. At a $B_1$ amplitude of 100%, the excitation bandwidth was between $-140$ Hz and 775 Hz for the CA-T2-Prep+FS and between $-66$ Hz and 601 Hz for the PA-T2-Prep.

As expected based on our pulse design, simulation results show that the choice of $\Delta \varphi$ affects the level of fat suppression (Figure 3A), whereas it does not affect the on-resonance signal. The optimal $\Delta \varphi$ for a given $T_2$-Prep duration ($d_1$) is found, the optimal $\Delta \varphi$ for another $T_2$-Prep duration ($d_2$) can be estimated as follows (derivation in Supporting Information):

$$\Delta \varphi_2 = \sin^{-1} \left( e^{\frac{d_2-d_1}{T_2^*}} \sin \left( \Delta \varphi_1 - \frac{\pi}{2} \right) \right) + \frac{\pi}{2} \tag{2}$$

### 3.2 Phantom studies

The minimum average fat signal occurred at $\Delta \varphi$, ranging from $105^\circ$ to $135^\circ$, depending on the $T_2$-Prep duration (Figure 3B). The choice of $\Delta \varphi$ only affects the level of fat suppression (Figure 3B), and it does not affect the on-resonance signal nor is it influenced by changes in $T_1$ or $T_2$ (Figure 3C-H).

The SNR of the blood, myocardium, and fat compartment using either the CA-T2-Prep, CA-T2-Prep+FS, WSA-T2-Prep+FS, and PA-T2-Prep relative to using no $T_2$-Prep show differences in fat suppression and $T_2$-weighting (Figure 4). Compared with the CA-T2-Prep, CA-T2-Prep+FS and WSA-T2-Prep+FS, the PA-T2-Prep reduced the lipid signal by 93.2%, 70.0%, and 66.1%, respectively. The blood and myocardium compartment SNR in PA-T2-Prep images were, respectively, 0.7% and 0.9% lower than in the CA-T2-Prep, 0.1% and 3.8% lower than in the CA-T2-Prep+FS, and 3.4% and 26.9% lower than in the WSA-T2-Prep+FS (Figure 4).

Compared with the CA-T2-Prep, CA-T2-Prep+FS and WSA-T2-Prep+FS, the PA-T2-Prep increased blood–myocardium CNR by −0.5%, 3.4%, and 35.3%, the blood–fat CNR by $-1399\%$, 22.5%, and 13.1%, and the myocardium–fat CNR by $-166\%$, 62.3%, and $-7.2\%$ (Figure 4).

Fat suppression using the PA-T2-Prep was improved in regions where field inhomogeneities hindered the capabilities of other methods (Figure 4, orange arrows). In contrast to the
CA-T₂-Prep+FS and PA-T₂-Prep, the WSA-T₂-Prep reduced blood and myocardium signals in regions where the B₀ imperfections increased the precession frequency by more than 200 Hz (Figure 4, green arrows).

3.3 | In vivo study

The PA-T₂-Prep visually increased the RCA vessel conspicuity compared with the CA-T₂-Prep+FS and the CA-T₂-Prep (Figure 5A). The PA-T₂-Prep reduced chest fat SNR to 7.2 ± 1.9 from 34 ± 10.8 (CA-T₂-Prep, P < .005) and 15.3 ± 5.2 (CA-T₂-Prep+FS, P < .05; Figure 5B). Epicardial fat SNR was reduced to 2.5 ± 0.8 (PA-T₂-Prep, P < .005) to 10 ± 2.5 (CA-T₂-Prep, P < .05) and 4.6 ± 1.9 (CA-T₂-Prep+FS, P < .05; Figure 5B). No significant differences between blood SNR and myocardium SNR was observed across the three different T₂-Prep approaches (Figure 5B), similar for the CNR between blood and myocardium (Figure 5C). The PA-T₂-Prep results reported the highest CNR between blood and epicardial fat, as well as between myocardium and epicardial fat (Figure 5C). Using the PA-T₂-Prep approach the RCA vessel sharpness was 40.2 ± 9.6%, compared with 21.0 ± 6.4% (CA-T₂-Prep, P < .005) and 33.6 ± 6.4% (CA-T₂-Prep+FS, P = .08; Figure 5D). The SAR increased by 1% using the PA-T₂-Prep compared with the two other methods.

4 | DISCUSSION

As shown by the Bloch equation simulations, the PA-T₂-Prep not only increases the fat saturation bandwidth, but also avoids water signal attenuation when its resonance frequency is shifted caused by field inhomogeneities. Additionally, the multiple rotations of the fat magnetization during the PA-T₂-Prep decrease the fat suppression
dependence on $B_1$ precision compared with the use of a single fat saturation pulse.

The WSA-T2-Prep\textsuperscript{10} achieved fat suppression by increasing the RF excitation angle of tip-down and tip-up pulses, which affects on-resonance magnetization and results in a loss of T$_2$ preparation. In comparison, the PA-T2-Prep rotates the magnetization 90° such that the effectiveness of the T$_2$ preparation is conserved, as confirmed by the measured SNR in the phantom and in vivo experiments. The water-fat–phase separation $\Delta \varphi$ influences the orientation of the fat magnetization.

FIGURE 4 A comparison of the different T$_2$-preparation (T$_2$-Prep) methods in the multicompartment “heart” phantom with corresponding $B_0$ and $B_1$ maps. Images acquired with no T$_2$-Prep, the conventional adiabatic T$_2$-Prep (CA-T$_2$-Prep) without and with fat saturation (FS), the water-selective adiabatic T$_2$-Prep (WSA-T$_2$-Prep) and the phaser adiabatic T$_2$-Prep (PA-T$_2$-Prep). Orange arrows indicate locations of improved fat suppression using the PA-T$_2$-Prep compared with other fat-suppression methods. Green arrows indicate positive off-resonance regions in which the water-selective method attenuated the signal that was not observed using the PA-T$_2$-Prep. The signal-to-noise ratio (SNR) and contrast-to-noise ratio (bottom panels) were computed for each technique. The blood and myocardium signals in the CA-T$_2$-Prep, CA-T$_2$-Prep+FS, and PA-T$_2$-Prep were similar. However, the 120° radiofrequency-excitation angle of WSA-T$_2$-Prep reduced the efficiency of the T$_2$ preparation, resulting in increased myocardium SNR. The fat signal was most reduced using the PA-T$_2$-Prep.
after the T2-Prep (Figure 1) and can be adjusted to adapt the module to different pulse sequences. For example, a balanced steady-state free-precession acquisition may require ramp-up pulses between the T2-Prep and the acquisition during which the fat signal may recover. In this case, $\Delta\phi$ may be increased to compensate for this additional recovery.

Similar to the CA-T2-Prep+FS, the WSA-T2-Prep, or a CHESS fat-saturation pulse, the PA-T2-Prep suppresses the fat once before the acquisition. After suppression, the fat magnetization will start to recover; consequentially, the effectiveness of the fat suppression is limited by the pulse sequence that follows. Although simulations provided a good initial estimate...
of the optimal $\Delta \varphi$, within an imaging sequence there are many parameters that affect the final signal weighting of fat. It is complex to simulate accurately; hence, there are small discrepancies in optimal $\Delta \varphi$ between simulations and experiments and the need for an experimental fine-tuning. Once $\Delta \varphi$ is found, it can be transferred to the in vivo setting provided that the same acquisition parameters are used, and that the phantom adequately represents the fat tissue of interest. Otherwise, an experimental fine-tuning of $\Delta \varphi$ is required in vivo. In sequences using Cartesian sampling trajectories, the PA-T$_2$-Prep may remove the need for additional fat-suppression modules and thus reduce SAR. However, it will be less suitable for acquisition schemes that do not allow for centric reordering of the phase-encoding planes of k-space, such as radial acquisitions. Although water-selective excitation approaches may be more suitable for fat suppression in radial whole-heart MRI at 3T, acquisitions requiring T$_2$-Prep modules may still benefit from the proposed approach because it offers an additional, tuneable range of fat suppression. Recently, T$_2$-Prep approaches designed for coronary artery imaging at 1.5T included outer volume-suppression strategies, with integrated spectral spatial sinc pulses for fat suppression. Their utility at 3T remains to be investigated, considering SAR may be higher and balanced steady-state free-precession less performant at higher field strengths; therefore, these methods were not compared in the current study. Another limitation includes the minimum duration of the PA-T$_2$-Prep module that the lengthy tip-down pulse imposes. The PA-T$_2$-Prep requires at least 29 ms compared with 21.7 ms for the CA-T$_2$-Prep, which may limit the range of available T$_2$ contrast, although this may be addressed using a shorter adiabatic refocusing pulse pair. It should be noted that the glycerol and olefinic fat resonances at 5.3 and 4.2 ppm fall within the excitation formant at higher field strengths; therefore, these methods would not be suppressed. Because these resonances constitute approximately 9% of the fat signal, this is not considered a significant disadvantage.

The proposed PA-T$_2$-Prep has a narrow excitation bandwidth that may result in on-resonance water being suppressed when $B_0$ conditions are not optimal. In the phantom experiments, a slight decrease of $<1\%$ (yet above the noise level) in blood and myocardium compartment signal was observed. Besides the narrow excitation bandwidth, this slight loss could also be explained by the excitation profile ripples of the phaser tip-down pulse caused by its finite duration. Nevertheless, strong effects of diminished on-resonance excitation were not observed.

A translation of the PA-T$_2$-Prep to different field strengths is possible; higher magnetic field strengths would require a decrease in the duration of the tip-down RF excitation pulse, and lower magnetic field strengths would require an increase. Furthermore, the principle of our PA-T$_2$-Prep design could be translated to any magnetization preparation module. Calibrations would need to be performed to find the optimal $\Delta \varphi$, but there are no theoretical barriers. For example, in T$_2$-mapping techniques that use incrementing T$_2$-Prep durations, it may reduce chemical shift artifacts that hinder cartilage delineation and quantification, and may obviate the use of additional fat-suppression modules. It could also be integrated with T$_1$- and T$_2$-mapping techniques. The approach may be extended to applications that use T$_2$-Prep modules combined with inversion recovery, as well as modules for motion-sensitized driven-equilibrium for blood signal suppression.

5 | CONCLUSION

In this study, a novel T$_2$-Prep approach was developed that provides a large spectral bandwidth of fat suppression. The technique is robust against $B_0$ and $B_1$ inhomogeneities, and preserves blood-myocardium SNR and CNR. As a result, the RCA vessel sharpness was increased compared with conventional approaches.

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DATA AVAILABILITY STATEMENT

All data and code are available upon request.

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**SUPPORTING INFORMATION**

Additional Supporting Information may be found online in the Supporting Information section.

**FIGURE S1** A comparison of the small flip angle approximation (left) and a Bloch simulation (right) of the RF pulse profile (Figure 1E). The difference between these two predictions are due to the violation of the assumption of small flip angle. Within the excitation bandwidth (±700 Hz), the ripples of the magnetization phase have an amplitude of approximately ±5°, but the effect of these might not be visible in practice since they are averaged out across the wide frequency range of fat. The phase behavior outside the excitation bandwidth can be neglected since the amplitude of the transversal magnetization is negligible.

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