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Feasibility of Ultra-high Simultaneous Multi-slice and In-plane Accelerations for Cardiac MRI Using Outer Volume Suppression and Leakage-Blocking Reconstruction

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Synopsis

Simultaneous multi-slice (SMS) imaging has gained increasing interest for enabling high scan-time acceleration at the cost of minimal loss in SNR. However, its applications in cardiac MRI have been limited, as the feasible acceleration is restricted by unfavorable coil geometry. In this study, we investigate the use of outer-volume suppression (OVS) in combination with CAIPIRINHA to promote dissimilarities among the multi-bands. We propose a time and SAR efficient multi-band scheme for OVS and apply these techniques with a leakage blocking reconstruction to increase the feasible acceleration in cardiac cine and perfusion imaging. Combining these techniques, we achieve clinical image quality with 5 fold SMS acceleration in Cine and 16-fold spatial-only acceleration in perfusion MRI.

Introduction

Simultaneous multi-slice/Multi-band (SMS/MB)¹ imaging has been introduced as a means for scan-time reduction, where the only loss in SNR is caused by coil-geometries. CAIPIRINHA² can be utilized to further reduce g-factors, using differential FOV shifts in each multi-band. However, applications in cardiac MRI have been limited due to coil geometry and presence of extra-cardiac tissue, such as chest and back fat, which leads to residual aliasing or leakage. This challenge is further exacerbated when using additional in-plane acceleration, as in most cardiac MRI scans.

In this study, we propose a fast outer-volume suppression (OVS) method to enable higher acceleration factors in cardiac imaging. We study cardiac cine and perfusion imaging, with interleaved repeated OVS, and a spatial-only (i.e. no assumptions about dynamic imaging) leakage-blocking reconstruction to enable very high acceleration rates.

Methods

Sequence and Reconstruction: Figure 1 depicts the sequence diagrams of the proposed methods for OVS-SMS/MB imaging. In both sequences, OVS is performed using the multi-band combination of two slab-selective saturation pulses. These are used to suppress signal from parallel slabs along phase-encoding dimension. OVS is performed in an interleaved manner to ensure consistent suppression throughout acquisition. In both sequences CAIPIRINHA is performed by differentially modifying the excitation pulse phase of each respective band in a cyclic fashion. Images were reconstructed using a leakage-blocking approach, called split slice-GRAPPA (Split-SG).

Imaging Experiments: In-vivo imaging was performed in healthy subjects at 3T (Siemens Magnetom Prisma). First-pass perfusion imaging was performed with injection of 0.05 mmol/kg gadobutrol (Gadovist) at 4 mL/s, followed by a 10-mL saline flush; and cine imaging was performed subsequently for time-efficient scanning. Cine parameters were: TR/TE/FA=4.3/2.1ms/12°, FOV/resolution=320x320/1.7x1.7mm², slice thickness=6mm, SMS/MB-factor=5 or 6, temporal resolution=41ms, breath-hold duration=15-17s. Further acceleration was assessed by retrospectively undersampling the MB=5 dataset by in-plane acceleration=2 while keeping ACS=24 reference lines. Perfusion parameters were: SMS/MB-factor=3, in-plane acceleration=4, and partial-Fourier=6/8 were utilized for an overall 16-fold acceleration. The imaging parameters were: FOV=360x360mm², resolution=1.7x1.7mm², slice-thickness=8mm, TR/TE/FA=2.9/1.7ms/12°, temporal resolution=110ms, saturation time=150ms.

Both sequences share the same OVS module with saturation slab=150mm (each side), 3.8ms asymmetric sinc, BWT=8, RF peak shift⁴ = 15%. MB-OVS preparation was interleaved between every 9 imaging pulses for perfusion and 8 for cine.

Results

The effects of OVS are depicted in Figure 2, which shows cine images without acceleration. Thorough saturation of the periphery along the phase-encoding direction is observed in all cardiac phases. Evaluation of signal intensity along phase-encoding direction shows a profound drop outside the region-of-interest, especially in the chest and back.

Figure 3 shows cine imaging with SMS/MB-factor=5. Without OVS (left), substantial signal leakage onto the heart prevents diagnostic information. Excellent image quality is obtained with OVS and Split-SG reconstruction (middle), resulting in artifact-free depiction of cardiac structures through all cardiac phases and slices. With the retrospective in-plane acceleration, for total 10-fold acceleration (right), proposed approach results in leakage-free images, although increased noise amplification is observed.

Figure 4 depicts SMS/MB-factor=6 cine images. Heavy leakage artifacts cover the heart without OVS (left). Proposed approach successfully eliminates leakage artifacts (right), depicting clear myocardium blood-interface. Residual contrast variation is observed, which might hamper image evaluation.

Perfusion images acquired with 16-fold spatial-only acceleration using the proposed approach are shown in Figure 5, with accurate representation of contrast dynamics across 9 slices per heart-beat. Despite low base-line SNR in the saturation-recovery sequence, no leakage artifacts are observed.

Discussion

We used OVS to reduce leakage from extra-cardiac tissues in cardiac SMS imaging. Compared to previous OVS pulses for cardiac MRI⁵⁻⁶, our scheme benefits from low SAR, especially important at 3T. This facilitates repeated application, ensuring homogeneous signal suppression throughout cardiac cycle. Furthermore, the brevity of the pulse minimizes disruption of the steady-state signal during the FLASH acquisition, mitigating potential artifacts that might otherwise occur.

As opposed to other methods for scan-time acceleration that are commonly used in cine MRI⁷ or perfusion⁸, proposed acquisition and reconstruction do not employ any interrelation in temporal dimension, precluding temporal blurring artifacts. Furthermore, no regularization is applied during the Split-SG reconstruction, circumventing risk of over-regularization, which may lead to patchy or blurry images.

Conclusion

The proposed MB-OVS technique with leakage-blocking reconstruction enables ultra-high scan time acceleration in cardiac MRI, without the need for regularization or exploiting temporal dependencies. MB-factors up to 5, and spatial only net-acceleration rates of 10-fold and 16-fold were achieved in cine and perfusion imaging, respectively, while maintaining diagnostic image quality.

Acknowledgements

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References

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Figures

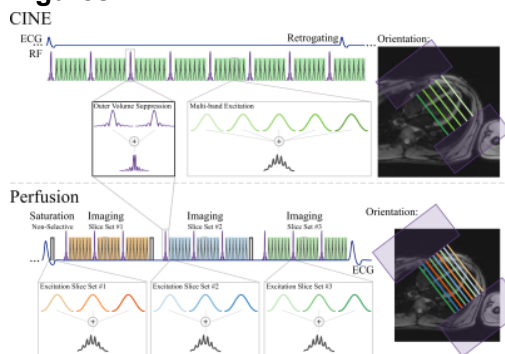


Figure 1: Sequence schemes for the proposed SMS-accelerated cine and perfusion sequence. Excitation pulses for FLASH imaging are obtained as the sum of 5 (cine) or 3 (perfusion) sinc-pulses at different center frequencies. OVS is achieved by interleaving these, with multi-band rest-slabs to simultaneously saturate two parallel slabs. Example orientation for short-axis imaging with phase-encoding along the anterior-posterior direction is illustrated on the right for both acquisitions.

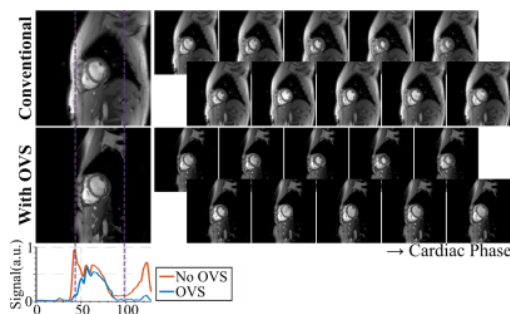


Figure 2: Illustration of the proposed outer-volume suppression (OVS) in cine imaging. Thorough signal depletion along the phase-encoding dimension (anterior-posterior), can be observed when applying the OVS module. Accordingly, signal intensity is substantially reduced outside the area of interest, especially near the back and the chest fat. The suppression is homogeneously maintained throughout all phases of the cardiac cycle.

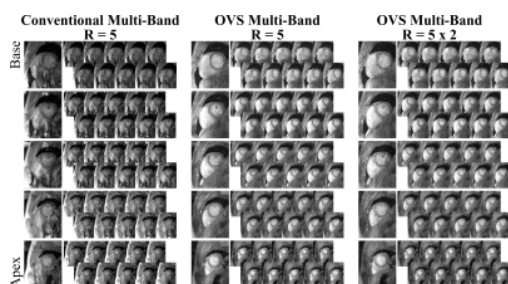


Figure 3: Example cine images acquired with 5-fold multi-band acceleration. Without the use of OVS severe leakage artifacts affect the region-of-interest, rendering non-diagnostic image quality (left). With OVS and leakage-blocking reconstruction (middle), no residual aliasing artifacts are observed, with clear blood-myocardial interface at the left-ventricle, suitable for quantitative evaluation. The proposed technique, maintains artifact free images when additionally applying two-fold in-plane acceleration, leading to 10-fold overall acceleration (right).

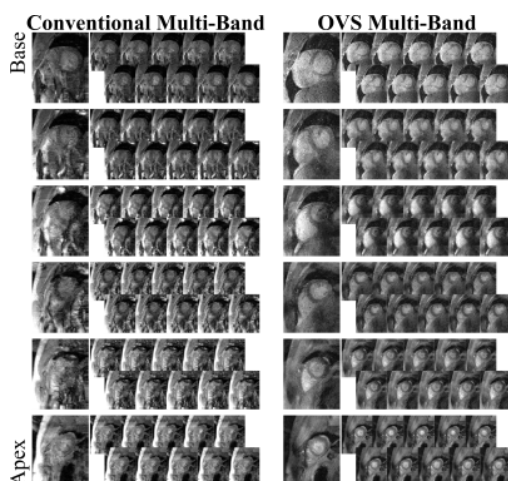


Figure 4: Example cine images acquired with 6-fold multi-band acceleration. The images acquired without OVS suffer from significant residual aliasing (left). These are mitigated with the proposed approach (right), which leads to images with clear delineation of the blood-myocardium interface.

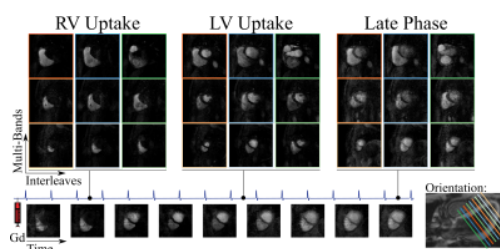


Figure 5: Example perfusion images acquired with the proposed OVS, using 3-fold multi-band acceleration, 4-fold in-plane acceleration and 6/8 partial Fourier imaging, for a total of 16-fold acceleration. This acceleration rate results in $1.7 \times 1.7 \text{ mm}^2$ spatial and 110ms temporal resolution, while offering whole heart coverage. Images across dynamics show high image quality during the contrast uptake in RV and LV, as well as during the later dynamics over the whole heart. Blood-pools are clearly delineated from the myocardium and no residual inter-slice leakage or blurring is apparent.