Optimized Parallel Imaging for Dynamic PC-MRI with Multi-Directional Velocity Encoding

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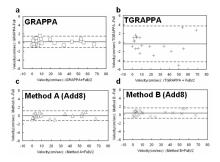
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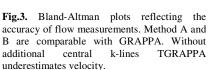
Introduction: Parallel imaging using GRAPPA has been used to accelerate image acquisition via undersampling of k-space in combination with properly-arranged RF coil arrays (1). In GRAPPA, AutoCalibrating Signals (ACS) are acquired for calculating the necessary reconstruction weights (WS). For dynamic phase-contrast (PC) MRI using GRAPPA, several ACS are typically acquired separately for each velocity direction and each cardiac phase. To further accelerate PC-MRI, we developed two methods, which calculated WS with reduced ACS data (2). In this study, we improve the image quality by retaining central part of k-space. The effects on flow velocity measurements were quantitatively investigated and compared to fully acquired data, standard GRAPPA (1), and TGRAPPA (time-interleaved sampling scheme in combination with GRAPPA) (3) reconstruction. In addition, the proposed algorithms were applied to tissue phase mapping data.

Methods and Materials: A full three-directional PC-MRI data set was acquired (3T Trio, Siemens, Germany, FOV=320x200mm, matrix=256x160, TR=40.8msec, VENC=1.5m/sec, 20 cardiac phases). k-lines were removed off-line to simulate GRAPPA with reduction factor (R) of 2, 3, and 4 and 32 central ACS lines (Fig.1(a)). To further accelerate data acquisition Method A computed WS from ACS of the reference image only (V0), which were applied to other three directionally encoded velocity data (Fig.1a). Method B estimated only the WS of V0 for the first cardiac phase (t1), which was then used to reconstruct all remaining images (Fig.1b). To improve the accuracy of flow measurements, images were also reconstructed with retaining 4 (Add4) or 8 (Add8) central k-lines. To evaluate different reconstruction strategies, blood flow velocities within the aorta (white circle in Fig.2a) of standard GRAPPA, TGRAPPA, and the proposed methods A and B were compared with the fully sampled data on a pixel-by-pixel basis. Images were reconstructed with R=[2,3,4] and [0,4,8] retained central k-lines. The new method (R=3, ACS=32, Add4) was also applied to tissue phase mapping data (matrix=256x96, TR=5.8ms, through-plane VENC=25 cm/sec, in-plane VENC=15cm/sec, 11 cardiac phases).

Results: Fig.2 shows magnitude images of fully sampled, standard GRAPPA, and the two proposed algorithms (R=2). Difference images demonstrated only subtle discrepancies for all methods. In Bland-Altman plots (Fig.3), Method A and B (Add8) were comparable with standard GRAPPA. In contrast, TGRAPPA underestimated flow velocity. For all investigated reconstruction algorithms Fig.4 shows the correlation coefficients (r²) for flow velocities, standard deviation (STD) for Bland-Altman plots, root mean square error (RMSE) for magnitude images, and % of time saving compared to GRAPPA. Methods A and B demonstrated good consistency (r² = 0.86~0.98) with fully sampled images (Fig.4(a)), and comparable STD (Fig.4(b)) and RMSE (Fig.4(c)) while providing additional reduction in scan time of 10%~17% (R=2) and 15%~28% (R=3) (Fig.4(d)). The application of Method A and B to tissue phase mapping data (Fig. 5) resulted in myocardium velocities comparable with standard GRAPPA and fully sampled data.

Discussion and Conclusions: The two proposed methods based on reduced ACS data acquisition can improve scan efficiency of dynamic PC-MRI while maintaining image quality and accuracy of measured flow velocities. The usage of retained central k-lines (Add4 and Add8) can improve the accuracy of flow measurements. For R=4, however, the images reconstructed with all reconstructed methods are deteriorated. Compared to TGRAPPA, Method A and Method B have proved to provide more accurate results (smaller STD) in evaluating flow velocity. For Add8, the proposed methods are comparable with standard GRAPPA. In contrast to k-t GRAPPA (4), the proposed methods can be implemented using the standard GRAPPA reconstruction. In conclusion, the two proposed reconstruction provide additional imaging acceleration of up to 25% for dynamic and multidirectional encoded acquisitions for R=2 and R=3.





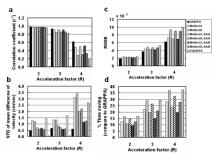


Fig.4. r² (a), STD (Bland-Altman plot) (b), and RMSE (c) for all methods for R=2,3,4 and added central k-lines. The results of Method A and B (with R=3, ACS=32, Add4 or Add 8) are comparable with GRAPPA.

References: 1. Griswold MA et al, 2002, MRM, 47(6):1202-1210. **2.** Peng HH et al, 2008, ESMRMB, No. 760. **3.** Beuer FA et al, 2005, MRM, 53(4):981-5. **4.** Huang F et al, 2005, MRM, 54(5):1172-84.

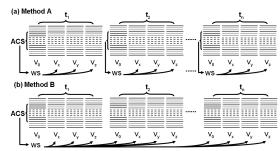


Fig.1. Method A (a) & B (b) with R=3. Solid line: acquired line. Dotted line: missing line. Dashed-line: retained central k-line. ACS: auto-calibrating signal. WS: coil weight estimation.

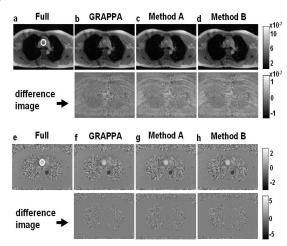


Fig.2. Magnitude images (a-d) and flow images (e-h) reconstructed with different strategies. The difference images (compare to fully sampled data) are shown in lower rows

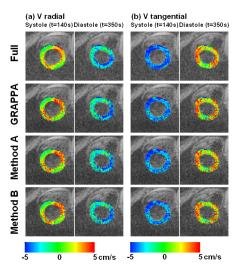


Fig.5. Myocardial motion velocities in radial (a) and circumferential (b) directions. Results of Methods A and B are comparable to standard GRAPPA and fully sampled data.