

A Clinically Applicable Scheme of MRI Trajectory Optimization for 3D Cartesian Acquisition

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Target Audience: MR researchers and clinical scientists working on fast acquisition.

Purpose: Random undersampling is an important component used with Parallel Imaging^{1, 2} (PI), Compressed Sensing³ (CS) and their combination (PI-CS)⁴ for fast acquisition. Optimized pseudo-random trajectory results in better reconstruction yet the computation complexity of optimizing undersampling trajectory prevented it from clinical application. Lately, we proposed an efficient scheme^{5, 6} for 1D random undersampling optimization using stochastic method and reference k -space data. Here we extended and improved the scheme to optimize the 2D Cartesian undersampling in 3D acquisition for PI and CS, by using Nonlinear Grappa Operator⁷ and Coherence⁸ based objective function. In-vivo experiments showed improved reconstruction using PI-CS.

Methods: The basic optimization scheme is similar to 1D undersampling optimization scheme^{5, 6} which optimizes the subset of k -space to be sampled using reference k -space from previous scans of similar positions and/or with different contrasts. The flowchart is shown at Figure 1. The main concept of the efficient scheme is: to replace computational-costly reconstruction steps in stochastic optimization iterations with efficient Pseudo-reconstruction^{5, 6} which only uses linear combination of pre-computed Parallel Imaging Operator results, and to use an objective function to approximate RMSE based on the Pseudo-reconstruction error. **To better optimize 2D trajectories, we chose to use a nonlinear GRAPPA operator⁷ (fig. 2) and a Coherence based objective function to better approximate the RMSE of PI-CS.**

1. A modified Simulated Annealing was implemented with a Gene-Algorithm based heuristic to optimize 2D sampling trajectories⁶.
2. A 3x3 Nonlinear GRAPPA Extrapolation operator⁷ was calibrated using the reference k -space to accurately extrapolate the neighbour samples. Since the kernel size is very small, nonlinear operator which maps the data to a higher dimension of virtual coils performs better than conventional GRAPPA. The extrapolation result was only computed once since they are not related to undersampling trajectories.
3. **Pseudo-Reconstruction:** For any arbitrary trajectory, un-acquired samples in k -space can be recovered using the pre-computed Nonlinear GRAPPA-operator results from sampled neighbour k -space data, which only needs efficient linear combination.
4. **Coherence based Objective Function** was used to compare two trajectories and update current optimal solution in iteration of stochastic optimization. Instead of error peak, we defined a new objective function to achieve more accurate prediction of the recovery performance using both PI and CS. In CS, incoherence of the sampling matrix D , which equals to the peak non-dc value in point spread function³, is the key for the convergence of l_1 regularized optimization³. Here we used the modified PI-recovery based point spread function (PSF) $psf(D) = \mathcal{F}^{-1}\hat{D} = \mathcal{F}^{-1}(D + E(D))$, in which \mathcal{F} for Fourier transform. \hat{D} is the pseudo-sampling operation resulted from pseudo-reconstruction results, which is a combination of D , sampling operation (1 for sampled points), and E , the normalized accuracy of pseudo-reconstruction ($E \approx 1$ if perfectly recovered). The objective function was modified accordingly: $f(D) = \|psf(D)\|_p, p \gg 1$.

Results: Two brain T1w/T2w datasets (FFE sequence/ 230x230mm²/256x256 matrix/16 slices) were acquired on a Philips 3T system (Philips Healthcare, Best, the Netherland) with an 8-channel head coil (Invivo Corporation, Gainesville, FL). The datasets were all fully sampled and retrospectively under-sampled. 2D random sampling trajectories were optimized and compared with VD Poisson Disk random trajectory⁴. The optimization scheme took about 30 seconds on Matlab on a 2.4GHz Quad CPU 64bit system. The T1w dataset was used for trajectory training, and the optimized trajectory was applied for reconstruction of another image. Figure 3 compared the L_1 -SPIRiT reconstruction using optimized and un-optimized trajectories. Figure 3 showed the decreasing of objective function in iterations in the stochastic optimization. Figure 4 showed the approximation of RMSE of different trajectories regressed from the objective function with/without using Nonlinear GRAPPA.

Discussion: The proposed method efficiently optimized the 2D Cartesian trajectories in 3D acquisition. To tackle the problem with high degree-of-freedom, nonlinear GRAPPA operator was used to accurately extrapolate data in k -space and pre-computation enabled fast pseudo-reconstruction given any trajectory in stochastic optimization. The modified Coherence based objective function better approximates the performance of CS after PI-based recovery to better predict RMSE of PI-CS reconstruction. Results showed the scheme was efficient and effective for 2D Cartesian trajectory optimization. This scheme can benefit fast 3D acquisition and Multi-contrast imaging⁵. This scheme is also adaptable for non-Cartesian sampling by gridding the Nonlinear GRAPPA of non-Cartesian samples to Cartesian points in pre-computation step.

Conclusion: We presented an efficient scheme for 2D Cartesian trajectory optimization in 3D acquisition which can be applied clinically.

References: [1] Pruessmann et.al. MRM 1999 [2] Griswold, et.al. MRM 2002 [3] Lustig et. al. MRM 2007 [4] Lustig et. al. MRM 2010 [5] Gong et.al. ISMRM 2013 [6] Gong et.al. Sedona 2013 [7] Chang, et.al. MRM 2012 [8] Donoho, et.al. IEEE TIT 2006

