

Combining parallel detection of proton spectroscopic imaging (PEPSI) measurements with a data-consistency constraint improves SNR

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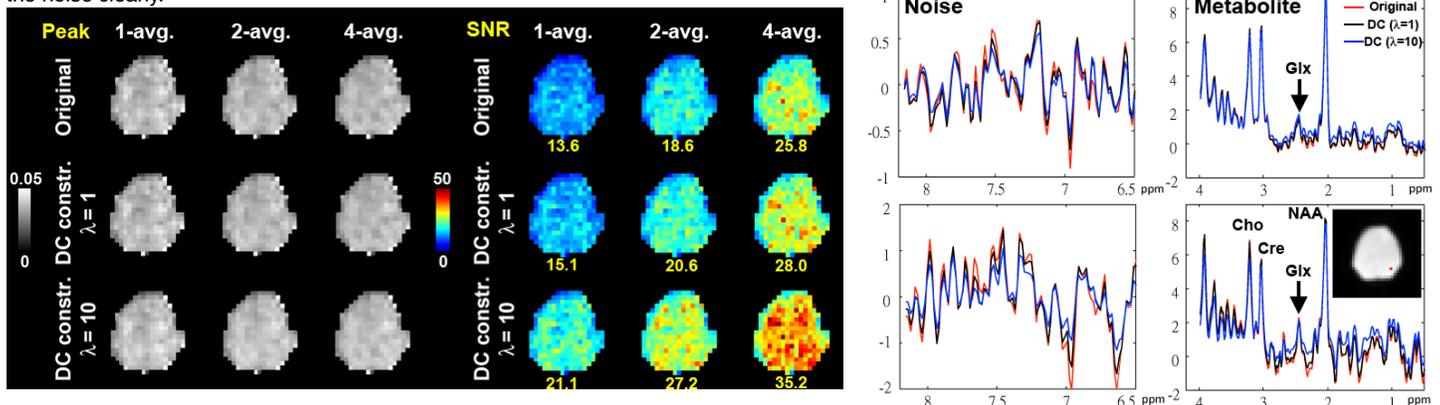
TARGET AUDIENCE Scientists interested in reducing noise in spectroscopic imaging data acquired from a coil array.

PURPOSE Fast MR spectroscopic imaging (MRSI) can be achieved using EPI¹ or spiral readouts². Yet one major challenge of MRSI is its relative low signal-to-noise ratio (SNR), which can be improved by using a surface coil array. Instead of utilizing the spatial sensitivity of different channels of a coil array to accelerate image encoding at the cost of the reduced SNR^{3,4}, here we propose to exploit this sensitivity information to enforce the *k*-space data-consistency (DC) in order to suppress noise and consequently to improve MRSI SNR. With *in vivo* experimental data at 3T using a 32-channel coil array, we found that the DC constraint can improve the SNR of PEPSI by approximately 40%.

METHODS Assuming the coil sensitivity profiles of a MRI detection array are distinct and spatially smooth, each chosen *k*-space data point of a coil can be expressed as the linear combination of the *k*-space data points from all coils at the vicinity of the chosen *k*-space data point. Mathematically, such DC relationship is $\mathbf{x} = \mathbf{G} \mathbf{x}$, where \mathbf{x} denotes the concatenation of *k*-space data from all coils and \mathbf{G} is a convolution kernel⁵. Practically, we first estimated the convolution kernel \mathbf{G} and then reconstructed images at each channel \mathbf{x} by minimizing the cost function $|\mathbf{S}\mathbf{x}-\mathbf{y}|^2+\lambda|\mathbf{G}\mathbf{x}-\mathbf{x}|^2$, where \mathbf{S} is an index matrix indicating the *k*-space coordinates of the acquired data, \mathbf{y} is the acquired *k*-space data across all channels, and λ is a regularization parameter.

PEPSI¹ was performed on healthy volunteers using a 3 T scanner (Tim Trio, SIEMENS Medical Solutions, Erlangen, Germany) and a 32-channel array. PEPSI data were acquired from a para-axial slice at the upper edge of the ventricles with TR 2 s and short TE (15 ms), using a 32x32 image matrix. Complete 8-slice outer volume suppression was applied along the perimeter of the brain to suppress lipid signal. Even- and odd-echo data were reconstructed separately using a non-water suppressed reference scan for automatic phasing and frequency shift correction. The total scan time for 1-, 2-, 4-average data were 64, 128, and 256 s. We used non-water suppressed image to estimate \mathbf{G} because it has a higher SNR. Then an iterative reconstruction method based on the conjugated gradient algorithm was used to minimize the cost for each time point of the water-suppressed scan. We chose $\lambda = 0$ (no DC constraint), 1, and 10 in this study. Finally, all images were reconstructed by sum-of-squares.

RESULTS Figure 1 shows the peak value and SNR maps of N-Acetyl-Acetate (NAA) using the standard deviation between 6.5 and 8 ppm as the noise estimate. The SNR in the brain parenchyma was slightly and significantly improved with DC constraint $\lambda = 1$ and 10 respectively. Comparing to the metabolite peak with $\lambda = 0$, the metabolite peaks were similar with $\lambda = 1$ and slightly decreased with $\lambda = 10$. However, the noise level was clearly suppressed in $\lambda = 1$ and further suppressed with $\lambda = 10$. Thus overall SNR (yellow numbers) is improved with $\lambda = 1$ and $\lambda = 10$. Similar improvement was observed for 1-, 2-, and 4-average data. Using the DC constraint with $\lambda = 10$, the SNR for 1- and 2-average data approximated the SNR for 2- and 4-average data without using the DC constraint. Figure 2 shows the spectra in metabolic range between 0.5 and 4 ppm as well as in noise range between 6.5 to 8 ppm. NAA, creatine (Cre), and choline (Cho) metabolite peaks were clearly identified in all spectra. Noise level was reduced by the DC constraint for both 1- and 4-average data. Interestingly, around 2.2 ppm, spectra show similar peaks, potentially from the glutamine/glutamate (Glx). Yet the DC constraint suppressed the noise clearly.



DISCUSSION We proposed a MRSI reconstruction algorithm enforcing the *k*-space data consistency among channels of a coil array such that the noise disturbing such a consistency is suppressed. Different from the optimal reconstruction of coil array images⁶, our method does not require explicit estimate of coil sensitivity maps, which in practice can be tedious. The results of using a DC constraint to significantly suppress the noise at the cost of marginally reducing the signal (metabolite peak) is similar to the advantage of using a prior to regularize parallel functional MRI reconstruction⁷. Yet our method does not need to explicitly specify the prior either. Choosing a larger regularization parameter ($\lambda = 1000$) can bias the reconstruction into the null space of \mathbf{G} (not shown). The proposed reconstruction algorithm can also incorporate image sparsity feature in order to further suppress noise. Our results suggest that the carefully optimized combination of MRSI data from a coil array can save around 50% of scanning time to obtain a similar image and spectra SNR.

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