

POSTER PRESENTATION

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Flip angle optimization for quantitative phase contrast MR imaging

Daniel B Ennis*, Matthew J Middione

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Objective

The overall objective of the work proposed herein is to optimize the flip angle in gradient-echo based PC-MRI methods for decreased measurement variability in quantitative blood velocity measurements of the common femoral arteries.

Background

PC-MRI is a noninvasive imaging technique used to measure the velocity of flowing blood in a particular blood vessel with flexible spatial and temporal resolution [1,2]. It is considered the clinical “gold standard” for quantification of blood flow. PC-MRI boasts a variety of established applications in quantifying cardiovascular function and hemodynamics. In particular, this technique offers clinicians with a means of measuring peak velocity, mean velocity, flow rate, and total flow throughout the vasculature of the human body. One of the challenges in PC-MRI is the wide range of variables that make up a given protocol requiring careful optimization of each imaging parameter in an effort to accomplish the goals of a given scan. Surprisingly, the current PC-MRI utilizes a default flip angle, which has not been optimized. A theoretical description of the signal acquired from flowing spins in spoiled gradient-echo pulse sequences has been previously developed [3]. These efforts indicate that there exists a flip angle that provides an optimized SNR for a given tissue, echo time, repetition time, slice thickness and velocity, which leads to increased reproducibility of quantitative PC-MRI measurements. This optimization cannot be performed empirically because of the number of parameters involved, hence mathematical and computation work is required.

Methods

All measurements were performed on a 1.5 Tesla system (Avanto; Siemens Medical Solutions, Erlangen, Germany) using a 16-channel surface coil. Blood velocity data in the common femoral arteries were obtained in 5 healthy consenting volunteers (4 male, 1 female, age \pm years). PC-MRI was performed without breath-hold using retrospective ECG triggering. Imaging parameters included 340 x 212.5 mm field of view, 256 x 160 matrix, 1.3 mm isotropic resolution, 8 mm slice thickness, 5.4/3.2 msec TR/TE, 32.4 msec temporal resolution, a range of flip angles for optimization (5,10,15,20,30,40,60,75 and 90°), 888 Hz/pixel acquisition bandwidth, three views per segment and a velocity encoding strength of 75 cm/sec (53 second scan time). The imaging was repeated 5 times in each volunteer.

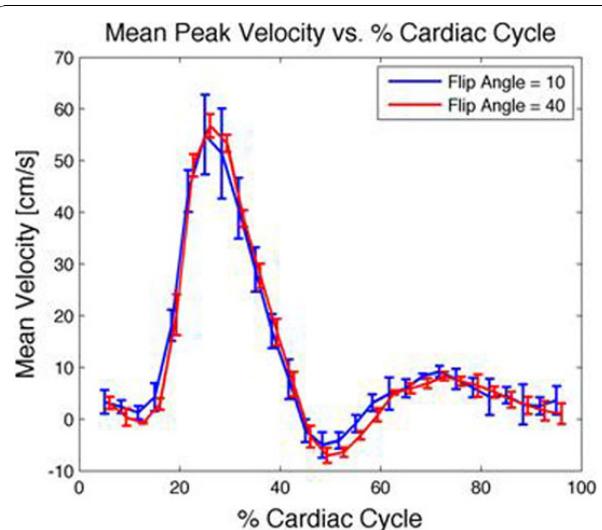


Figure 1

University of California, Los Angeles, Los Angeles, CA, USA

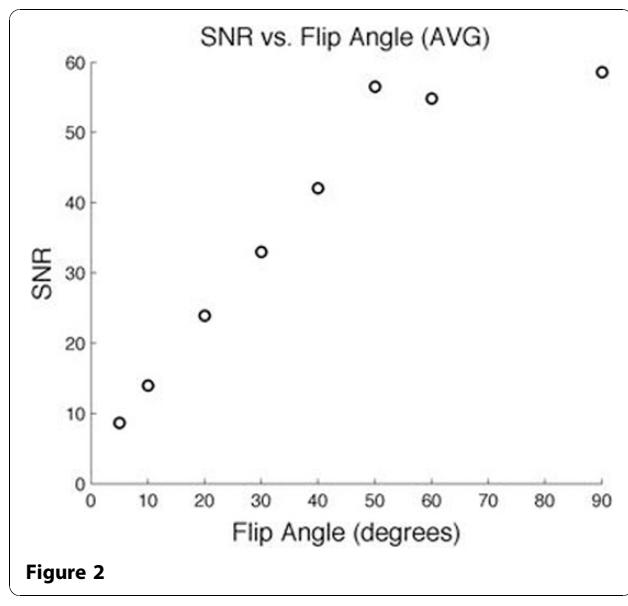


Figure 2

Results

The results indicate that there exists an optimal flip angle (Figure 1) in which the SNR is optimized for a given experiment. The use of this flip angle yields decreased variability in quantitative blood flow measurements compared to other non-optimized flip angles (Figure 2). From these results it can be concluded that careful selection of the flip angle is required. Flip angles for flow quantification are not sequence dependent; rather they are velocity dependent for a given protocol and simply selecting a default flip angle for any given study will not be sufficient to yield the best results.

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References

1. Pelc NJ: *Magn Reson Imaging* 1995, **3**:413-424.
2. J Lotz J, et al: *Radiographics* 2002, **22**:651-671.
3. Gao JH, et al: *Medical Physics* 1988, **15**:809-814.

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