# Parallel Magnetic Resonance Imaging (pMRI): Implementations, Problems and Future

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## Overview

- Quick Review
- Implementations of Parallel Imaging
- QA problems in Parallel Imaging
- QA solutions in Parallel Imaging
- Advanced Parallel Imaging and other reconstruction techniques





## Parallel Imaging: What is it good for?

Faster acquisition time

- Can be added to a majority of MR Protocols
- Complementary to other acceleration methods
- Cardiac Imaging
- Perfusion and Diffusion Imaging





# Parallel Imaging: How does it work?

Coil sensitivity profile found before or during acquisition

Multiple phased array coils acquire pieces of k-space

Pieces put together like a puzzle

Aliasing occurs

Coil sensitivity profile used to un-alias image







Image taken from: Larkman DJ, Nunes RG. Parallel magnetic resonance imaging. Phys Med Biol 2007 Apr;52(7):R15-55.





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Image taken from: Larkman DJ, Nunes RG. Parallel magnetic resonance imaging. Phys Med Biol 2007 Apr;52(7):R15-55.







Image taken from: Larkman DJ, Nunes RG. Parallel magnetic resonance imaging. Phys Med Biol 2007 Apr;52(7):R15-55.





Can provide faster acquisition times

- R- the reduction factor is the factor that the acquisition time is reduced by
- Trade-off is **reduced** signal to noise ratio **(SNR)** and an increase in residual aliasing artifacts

Typical R values are 2, 3, or 4



Normal

R=2



R=3

R=4



Artifacts with increased R for an FSE sequence with Parallel Imaging





# pMRI Acronyms by manufacturer

Acronym	Manufacturer	<b>Reconstruction Method</b>	Calibration Method
GRAPPA	Siemens	k-space	Auto
mSENSE	Siemens	image space	Auto
SENSE	Philips	image space	Pre-scan
ASSET	GE	image space	Pre-scan
ARC	GE	k-space	Pre-scan
SPEEDER	Toshiba	image space	Pre-scan







Introduced by Pruessman as a metric to indicate the decrease in SNR

Reduction achieved with coil sensitivity information

- Results in spatially varying noise and thus SNR
- g-factor accounts for this variation

g-factor varies with spatial position

- Useful images typically have a g between 1 and 2
- Difficult for clinical diagnostic physicist to obtain
- Requires knowledge of coil sensitivities

$$g = \frac{SNR_{R=1}}{SNR_R\sqrt{R}} \ge 1$$







Accurate SNR measurements difficult to obtain with Parallel Imaging implemented

Difficult to compare image quality across protocols and platforms

Most difficult to measure noise because it varies from pixel to pixel

$$SNR_R = \frac{SNR_{R=1}}{g\sqrt{R}}$$





### NEMA method 1: Image subtraction

- Signal: 80% average signal ROI
- Noise: 80% SD ROI of subtracted images





Method N2 Noise







Method N4 Noise



Method ACR Noise





Method N1 Noise





# NEMA method 2: No signal image

- Signal: 80% ROI
- Noise: 80% SD ROI of no signal image







# NEMA method 4: SD of background

- Signal: 80% ROI
- Noise: SD of 1000 pixels from background/0.66







ACR method: SD of smaller background portion

- Signal: 80% ROI
- Noise: SD of 50 pixels from background portion of image







# Sequences Compared

### Turbo Spin Echo (TSE)

• T<sub>2</sub> weighted images

Echo Planar Imaging (EPI)

• Functional MRI studies

Balanced Steady State Free Precession (TruFISP)

• Cardiac imaging





# Data Collection & Analysis

Each sequence was taken with two methods of auto calibrated parallel imaging at R=2,3 and 4

- mSENSE: image based recon
- GRAPPA: k-space based recon

Three acquisitions per protocol

- 2 for image subtraction
- 1 with RF voltage set to 0 V, for no signal method

Each SNR method implemented

- g-factor calculated for each method
- Best method should maintain g-factor>1 for R=2,3 and 4

$$g = \frac{SNR_{R=1}}{SNR_R \sqrt{R}} \ge 1$$



## Results

g-factor  $\geq 1$ 

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- Average of all sequences
  - NEMA 1: 2.01 ± 0.70
  - NEMA 2: 0.64 ± 0.22
  - NEMA 4: 0.81 ± 0.31
  - ACR: 0.64 ± 0.22

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	GRAPPA								
R	M100	N1	N2	N4	ACR				
2	$0.99 \pm 0.01$	$1.08 \pm 0.02$	$0.58 \pm 0$	$0.59\pm0.02$	$0.58 \pm 0.04$				
3	$1.15\pm0.01$	$1.31 \pm 0.02$	$0.44 \pm 0$	$0.4 \pm 0.01$	$0.38 \pm 0.03$				
4	$1.76 \pm 0.03$	$2.06 \pm 0.03$	$0.35 \pm 0$	$0.38\pm0.01$	$0.35 \pm 0.02$				
mSENSE									
R	M100	N1	N2	N4	ACR				
2	$1.04 \pm 0.12$	$1.13 \pm 0.19$	$0.81 \pm 0.05$	$0.9 \pm 0.09$	$0.89 \pm 0.07$				
3	$1.24 \pm 0.03$	$1.4 \pm 0.13$	$0.83\pm0.06$	$0.9 \pm 0.08$	$0.9\pm0.08$				
4	$1.92\pm0.04$	$2.28\pm0.35$	$0.88\pm0.11$	$0.93 \pm 0.13$	$0.92\pm0.09$				

Data taken from: Goerner FL, et al. Signal-to-noise ratio in parallel imaging MRI. Med Phys 2011 Sept;38(9)

## Conclusions

- **NEMA 1**: is the only method maintaining g>1
  - g=1.61 ± 0.62
- The ACR method consistently results in g<1
  - $g=0.44 \pm 0.31$



Recommendation: To compare SNR protocols using parallel imaging, the image subtraction method should be used





# Uniformity

### NEMA method 1 (UN1):

Peak deviation non-uniformity

NEMA method 2 (UN2):

Gray Scale Uniformity Map

#### NEMA method 3 (UTTT):

• Tic Tac Toe Method

ACR Method (UACR):

• Percent Image Uniformity

NAAD (UNAAD):

• Normalized Absolute Average Deviation





# Uniformity: UACR

UACR: Percent Image Uniformity

- Two ROI's encompassing 0.15% of the phantom volume (S<sub>max</sub> & S<sub>min</sub>)
- S<sub>max</sub>- area of greatest signal intensity
- S<sub>min</sub>- area of lowest signal intensity

$$UACR = 100 \left\{ 1 - \frac{(S_{\max} - S_{\min})}{(S_{\max} + S_{\min})} \right\}$$

Higher number indicates greater uniformity





# Uniformity: UACR



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# Uniformity: UN1









# Uniformity

### UN2: Gray Scale Uniformity

- Take the mean (m) from a 75% ROI
- Reassign pixel values (pv) according to difference from mean
  - i. -10%<pv<10% neutral
  - ii. 10%<pv<20% next brighter grey level
  - iii. -20%<pv<-10% next darker grey level
  - iv. pv>20% white
  - v. pv<-20% black





# Uniformity: UN2

- Provides visual uniformity map
- For numerical comparison
- Group number found by taking total number of pixels in that group and dividing by total number of pixels

Group 1

- i. 10%<pv<20%
- ii. -20%<pv<-10%

- Group 2
- iii. pv>20%
- iv. pv<-20%

### $UN2 = 100(1 - (0.5 \cdot Group_1 + Group_2))$





# Uniformity

### UNAAD

• Take a 75% ROI and find the average pixel value  $(\overline{Y})$ 

$$UNAAD = 100 \left( 1 - \frac{1}{N \cdot \overline{Y}} \sum_{i=1}^{N} \left| Y_i - \overline{Y} \right| \right)$$

 Where Y<sub>i</sub> is individual pixel value and N is the total number of pixels





# Uniformity

### **UTTT: Tic Tac Toe Method**

- S<sub>18</sub>- mean of 75% ROI
- 17 small 7x7 pixel ROI's
  - 9 in a tic-tac-toe pattern
  - 4 in the corners of the image
  - 4 in the middle edge of each side  $UTTT = 100 \times_{\text{Q}}^{\hat{\mathcal{Q}}} \frac{\stackrel{17}{\text{a}}}{\underset{\text{Q}}{\overset{17}{|S_n - S_{18}|}} \stackrel{0}{\underset{\text{M}}{\overset{\cdot}{|S_n + S_{18}|}} \stackrel{\cdot}{\underset{\text{M}}{\overset{\cdot}{\div}}}{\frac{17}{|S_n + S_{18}|} \stackrel{\cdot}{\underset{\text{M}}{\overset{\cdot}{\div}}}$
- S<sub>n</sub>- mean of small ROI
- n- ROI number

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## MRI Phantom

Phantom: Soccer ball provided by AAPM TG#118

- 19 cm outer diameter, 16.6 cm inner diameter
- filled with 5.45 g NaCl (99.99% pure) and 5.29 mL of Magnevist per 1 L distilled water
- Total volume: 2415 mL

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# Materials and Methods

#### **Pulse Sequences**

- Echo planar imaging (EPI)
- Fast Low Angle SHot (FLASH)
- True Fast Imaging with Steady-state Precession (Tru-FISP)
- Turbo Spin Echo (TSE)





## Variables

Two methods of reconstruction

- GRAPPA- k-space based
- mSENSE- image space based
- Varied R-values: 2,3,4

Varied phase encode:

Axial: AP and RL





## **MRI** Protocols

FOV=220 mm	Sequence	TR	TE	BW
1 mm slice gap		(ms)	(ms)	Hz/pixel
5 mm slice thickness	EPI	1840	187	752
	FLASH	175	4	240
256x256 matrix size	Tru-FISP	6.88	3.44	244
5 slices	TSE	1200	76	122







Linear fits for R-value vs. Uniformity

- Average slopes
- Two way ANOVA- R-value vs.
  - Reconstruction method
  - Pulse sequence
  - PE direction





## Noise Propagation Artifact

Decrease in uniformity with increasing R-value

#### Increase in noise propagation with mSENSE No PPI R=2 R=3



#### 3<sup>rd</sup> slice of FLASH sequence



**Image taken from: Goerner FL, et al.** A comparison of five standard methods for evaluating image intensity uniformity in partially parallel imaging MRI. **Med Phys 2013 Aug;40(8)** 

R=4

# g-factor map changes

With increase in R Decrease in uniformity



Images from: Breuer FA, et. al. "General Formulation for Quantitative G-factor Calculation in GRAPPA Reconstructions" MRM (2009) 62:739-746





## g-Factor maps

a-mSENSE (image space)

b- GRAPPA (k-space)

Arguably worse uniformity with mSENSE



Images from: Breuer FA, et. al. "General Formulation for Quantitative G-factor Calculation in GRAPPA Reconstructions" MRM (2009) 62:739-746





2

0

## UN1: Average Slope -4.0 $\pm$ 4.2

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## UN2: Average Slope $-1.03 \pm 1.4$



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## UACR: Average Slope $-0.50 \pm 0.82$





## UNAAD: Average Slope $1.02 \pm 1.8$



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## UTT: Average Slope: $0.004 \pm 0.2$





## Conclusions

UN1 and UN2 were more likely to have negative slopes (-4.0 and -1.2)

UN1 only uses two pixels and is sensitive to SNR

UN2 is difficult to measure clinically

There isn't really a good Uniformity measurement to characterize multi-channel coils and parallel imaging protocols.





# Advanced/Upcoming Techniques

Parallel imaging in 2 directions

CAIPIRINHA

**Compressed Sensing** 





# Parallel imaging 2 Directions

Current method- Reduce number of PE steps



**Frequency Encode** 

Frequency Encode





# Parallel imaging 2 Directions

Current method- Reduce number of PE steps



# 2 direction Parallel imaging

If R=2x2

Every other Phase Encode line is eliminated

Every other Slice Encode line is eliminated

Acquisition time reduced by 4x

Experimental techniques involve other trajectories not parallel to Phase or Slice encoding directions.





# CAIPIRINHA

First seen in 1918 in Sao Paulo Brazil

Now the National Drink of Brazil

Ingredients:

50 ml Cachaca

1/2 Lime (cut into four wedges)

2 teaspoons refined sugar



Short for: Controlled aliasing in parallel imaging results in higher acceleration





## CAIPIRINHA: The Quest



## CAIPIRINHA

Controlled aliasing in parallel imaging results in higher acceleration

- Multiple slices excited at once
- Phase encode is shifted with respect to other slices
- Creates a shift in aliasing artifact
- Potentially results in increased SNR, more R-values, fewer artifacts





# Multiple Slice Excitation







# Clinical Example

**3D VIBE Liver** 

Without pMRI

Acq Time= 43 seconds

With R=4

Acq Time = 17 seconds

With R=2x2

Acq Time = 11.4 seconds



CAIPI R=2x2

No pMRI





# **Compressed Sensing**

Similar concept to Parallel Imaging

Take less data

Different from Parallel imaging

Try to figure out what data you don't need and don't acquire it

JPEG images are compressed around 14x because of a lot of the information is similar. Compressed sensing works similarly.

Requires a lot of processing power!





# **Compressed Sensing**

illustrates 6 sequential, timeresolved MIP acquisitions during a contrast enhanced MR angiogram also acquired at 3T using a 32channel head coil.

Courtesy of Mark Griswold, Cleveland, OH







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