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PARIETAL

University of British Columbia Vancouver, Canada



THE UNIVERSITY OF BRITISH COLUMBIA Retrospective & Prospective SPARKLING Trajectories for Accelerated 2D Anatomical Imaging at 7T Using Compressed Sensing

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IEEE SPS Lecture Dec, 12 2016 DE LA RECHERCHE À L'INDUSTRI



## **INTRODUCTION: HIGH RESOLUTION IMAGING**



2D T2\*w axial, 7T scanner 120 x 120 x 600 µm<sup>3</sup> Matrix size: 1690 x 1744 21 slices, 2 averages 32-channel receiver coil, Motion correction, R=1

Acquisition Time of 50 minutes! *How can we* 

accelerate this?



## **INTRODUCTION: STANDARD MRI ACQUISITION**

Collect Fourier samples over a Cartesian grid



Idea: collect less Fourier samples to reduce acquisition time



## **INTRODUCTION: NAIVE PARTIAL FOURIER**





## **MRI ACQUISITION ALONG TRAJECTORIES**



## Mathematical modelling:

Let  $s : [0, T] \to \mathbb{R}^d$ , (d = 2, 3) denote the sampling curve. We have:

$$s(t) = s(0) + \gamma \int_0^t g(\tau) d\tau \operatorname{avec} g = (g_x, g_y).$$



Figure : Spiral imaging: Pulse sequence (Left) and corresponding sampling trajectory (Right).



K-space location is proportional to accumulated area under gradient waveforms



The gradient encoding g should satisfy: The g field is called gradient encoding, it should satisfy:

- $\|g\|_{\infty} \leq G_{\max}$ : bounded gradient magnitude, (eg, 70 mT.m<sup>-1</sup>).
- $\|\dot{g}\|_{\infty} \leq S_{\max}$ : bounded slew rate, (eg, 300 T.m<sup>-1</sup>.s<sup>-1</sup>).

## Admissible sampling curves

An admissible sampling curve in MRI is a curve belonging to the set:

$$\mathcal{S}_{\mathsf{MRI}} = \left\{ \boldsymbol{s} \in \left( \mathcal{C}^2([0, \, T]) \right)^d, \| \dot{\boldsymbol{s}} \|_{\infty} \leqslant \alpha = \gamma \boldsymbol{G}_{\mathsf{max}}, \, \| \ddot{\boldsymbol{s}} \|_{\infty} \leqslant \beta = \gamma \boldsymbol{S}_{\mathsf{max}} \right\}$$

Similar to driving a car on the Fourier plane.



Can we reduce the acquisition time by measuring fewer samples and still be able to reconstruct nice images?





- Part I: From CS to pointwise Variable Density Sampling
- Part II: Curve-based Variable Density Sampling
- Part III: Projection on measure sets
- Part IV: MRI simulation results
- Part V: Retrospective & Prospective SPARKLING at 7 Tesla

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# Part I: From Compressed Sensing to pointwise Variable Density Sampling

## **COMPRESSED SENSING RECIPE**

Data is sparse, compressible, redundant... Sense the compressed information directly!



Michael Lustig, http://www.eecs.berkeley.edu/~mlustig/CS.html



Donoho, Tao, Romberg, Candes

[Candes et al, IEEE IT 2006] [Donoho, IEEE IT 2006]



## Sparsity/Compressibility

#### Sparse

(spärs), *adj.* spars•er, spars•est.
1. Thinly scattered or distributed; not thic k or dense.
2. Scanty; meager.

(http://www.thefreedictionary.com/sparse)

#### Angiography image... is sparse



(www.healthcare.siemens)

![](_page_12_Picture_1.jpeg)

## WHAT SPARSITY AND COMPRESSIBILITY MEAN?

## Sparsity/Compressibility

#### Sparse

Compressible

(spärs), *adj.* spars•er, spars•est.
1. Thinly scattered or distributed; not thic k or dense.
2. Scanty; meager.

(http://www.thefreedictionary.com/sparse)

1. There exists a basis where the representation has just a few large

2. Compressible signals are well

coefficients and many small coefficients.

approximated by sparse representations

![](_page_12_Picture_7.jpeg)

is not sparse... ... but compressible!

![](_page_12_Picture_9.jpeg)

#### Wavelet Represensation... is sparse!

![](_page_12_Picture_11.jpeg)

3 levels of decomposition

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## **COMPRESSED SENSING THEORY (1/4)**

Compressed sensing theory:

- x is sparse in a given basis (e.g. wavelets):  $x = \Psi z$ , where  $z \in \mathbb{C}^n$  is s-sparse.
- Acquisition matrix: A = F<sup>\*</sup>Ψ.

Let  $\Gamma \subseteq \{1, \dots, n\}$  and  $A_{\Gamma} = (a_i^*)_{i \in \Gamma}$ . We acquire a measurement vector:

$$y = A_{\Gamma} z.$$

![](_page_13_Figure_7.jpeg)

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## **COMPRESSED SENSING THEORY (2/4)**

Compressed sensing theory:

- x is sparse in a given basis (e.g. wavelets):  $x = \Psi z$ , where  $z \in \mathbb{C}^n$  is s-sparse.
- Acquisition matrix:  $A = F^* \Psi$ .

Let  $\Gamma \subseteq \{1, \dots, n\}$  and  $A_{\Gamma} = (a_i^*)_{i \in \Gamma}$ . We acquire a measurement vector:

$$y = A_{\Gamma} z.$$

![](_page_14_Figure_7.jpeg)

 $\ell_1$  reconstruction (promoting sparsity)

$$\min_{\boldsymbol{z}\in\mathbb{C}^n, \boldsymbol{A}_{\Gamma}\boldsymbol{z}=\boldsymbol{y}}\|\boldsymbol{z}\|_1.$$

Compressed sensing theory:

- x is sparse in a given basis (e.g. wavelets):  $x = \Psi z$ , where  $z \in \mathbb{C}^n$  is s-sparse.
- Acquisition matrix:  $A = F^* \Psi$ .

Let  $\Gamma \subseteq \{1, \dots, n\}$  and  $A_{\Gamma} = (a_i^*)_{i \in \Gamma}$ . We acquire a measurement vector:

$$y = A_{\Gamma} z.$$

![](_page_15_Figure_7.jpeg)

or in case of noise (synthesis formulation):

$$\widehat{\boldsymbol{z}} = \operatorname*{Arg\,min}_{\boldsymbol{z} \in \mathbb{C}^n} \|\boldsymbol{y} - \boldsymbol{A}_{\Gamma} \boldsymbol{z}\|_2^2 + \lambda \|\boldsymbol{z}\|_1 \quad \text{(FISTA algorithm)}_{| \text{ page 16}}$$

## A first CS theorem [Candès and Plan, 2011]

#### Theorem

Construct  $\Gamma$  by uniform and i.i.d. drawing the lines of A. Let x be a sparse vector, containing s non-zero entries. Assume that:

$$m \ge C \cdot s \cdot \left( n \cdot \max_{1 \le k \le n} \|a_k\|_{\infty}^2 \right) \cdot \log\left(\frac{n}{\eta}\right)$$
(1)

where C is a universal constant. Then, with probability  $1 - \eta$ , x is the unique solution of:

$$\min_{\boldsymbol{z}\in\mathbb{C}^n,\boldsymbol{A}_{\boldsymbol{\Gamma}}\boldsymbol{z}=\boldsymbol{y}}\|\boldsymbol{z}\|_1.$$

In MRI,  $\max_{1 \leq k \leq n} \|a_k\|_{\infty}^2 = O(1)$ , hence  $m \gg n$ .

This is called the coherence barrier

![](_page_17_Picture_0.jpeg)

## A first CS theorem [Candès and Plan, 2011]

#### Theorem

Construct  $\Gamma$  by uniform and i.i.d. drawing the lines of A. Let x be a sparse vector, containing s non-zero entries. Assume that:

$$m \ge C \cdot s \cdot \left( n \cdot \max_{1 \le k \le n} \|a_k\|_{\infty}^2 \right) \cdot \log\left(\frac{n}{\eta}\right)$$
(1)

where C is a universal constant. Then, with probability  $1 - \eta$ , x is the unique solution of:

![](_page_17_Picture_7.jpeg)

![](_page_18_Picture_0.jpeg)

## Sample low frequencies more often

![](_page_18_Picture_3.jpeg)

(Lustig et al. 2007, Puy et al. 2011, Krahmer & Ward, 2012)

Breaking the coherence barrier: A new theory for Compressed Sensing Asymptotic incoherence, asymptotic sparsity and multi-level sampling [Adcock et al. 2013; Roman et al, 2014]

## Theorem [Chauffert et al., 2013]

Let x be an arbitrary *s*-sparse vector. Let  $(J_k)_{k \in \{1,...,m\}}$  denote a sequence of i.i.d. random variables taking value  $i \in \{1,...,n\}$  with probability  $p_i$ . Generate a random set  $\Gamma = \{J_1, \ldots, J_m\}$  and measure  $y = A_{\Gamma}x$ . Take  $\eta \in ]0, 1[$  and assume that:

$$m \ge C \cdot s \cdot \max_{k \in \{1, \dots, n\}} \frac{\|a_k\|_{\infty}^2}{p_k} \ln\left(\frac{n}{\eta}\right)$$

where C is a universal constant. Then with probability  $1 - \eta$  vector x is the unique solution of the following problem:

$$\min_{\boldsymbol{z}\in\mathbb{C}^n, A_{\mathsf{\Gamma}}\boldsymbol{z}=\boldsymbol{y}}\|\boldsymbol{z}\|_1.$$

Optimal distribution  $\pi_k \propto ||a_k||_{\infty}^2$ . Coherence is now  $\max_{k \in \{1,...,n\}} \frac{||a_k||_{\infty}^2}{p_k} = \sum_k ||a_k||_{\infty}^2 = O(\log(n))$  in MRI.

![](_page_20_Picture_1.jpeg)

#### Illustration of optimal sampling strategy for $H = F^* \Psi$ (MRI)

![](_page_20_Figure_3.jpeg)

Shannon Wavelets

#### Example of sampling pattern obtained in 2D :

![](_page_20_Figure_6.jpeg)

![](_page_21_Picture_1.jpeg)

![](_page_21_Figure_2.jpeg)

CS-MRI is sub-optimal! [Lustig et al., 2007]

## ➤CS not used to its full potential:

- Hindered randomness
- Variable density sampling not fulfilled in 3D
- K-space oversampled in one direction
- Undersampling factor generally limited to:  $R \leq 10$

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![](_page_22_Picture_1.jpeg)

## Part II: Curve-based Variable Density Sampling

[Chauffert et al, SIAM IS 2014, Chauffert et al, IEEE TMI]

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• CS must comply with MR hardware constraints

$$k(t) = k(0) + \gamma \int_0^t G(u) du$$
  

$$G < G_{max} \approx 50 \, mTm^{-1}$$
  

$$\dot{G} < \dot{G}_{max} \approx 333 \, mTm^{-1}s^{-1}$$

Regular trajectories

![](_page_23_Figure_5.jpeg)

Lustig et al. 2008

- Easy implementation: undersampling standard MR trajectories!
  - Radial for cardiac cine MR imaging (Winkelmann et al. 2007)
  - Spiral or noisy spirals (Lustig et al. 2005)
  - Poisson disk sampling (Vasanawala et al. 2011)

![](_page_24_Picture_0.jpeg)

## MATHEMATICAL FORMULATION OF VDS

## Pushforward measure - illustration

![](_page_24_Figure_3.jpeg)

![](_page_25_Picture_0.jpeg)

## MATHEMATICAL FORMULATION OF VDS

## Pushforward measure - illustration

![](_page_25_Picture_3.jpeg)

![](_page_26_Picture_0.jpeg)

## MATHEMATICAL FORMULATION OF VDS

### Pushforward measure - illustration

![](_page_26_Picture_3.jpeg)

 $\nu(B) = s_* \lambda_T(B) = \lambda_T(s^{-1}(B))$ 

 $\lambda_T$  is the (normalized) Lebesgue measure.

![](_page_27_Picture_0.jpeg)

## VDS - DEFINITIONS

#### Pushforward measure

Let  $\Omega = [0, 1]^d$ , where d = 2 or 3 denote the space dimension. We equip  $\Omega$  with the Borel algebra  $\mathcal{B}$ . Let  $(X, \Sigma)$  be a measurable space and  $s : X \to \Omega$  be a measurable mapping.  $\mu : X \to [0; +\infty[$  denote a measure. The *pushforward measure*  $\nu$  of  $\mu$  is defined by:

$$\nu(B) = s_*\mu(B) = \mu(s^{-1}(B)), \quad \forall B \in \mathcal{B}$$

### Ex. 1: Measures supported by curves

#### Ex. 2: Atomic measures

 $s: \{1, \ldots, m\} \to \Omega$ , where  $s(i) = p_i$  denotes the *i*-th point. Set  $\mu$  as the *counting* measure defined for any set  $I \subseteq \{1, \ldots, m\}$  by  $\mu(I) = \frac{|I|}{m}$ . Then  $\nu$  is defined by

$$\nu = \frac{1}{m} \sum_{i=1}^{m} \delta_{p_i}.$$

![](_page_28_Picture_0.jpeg)

#### Weak convergence [Chauffert et al, SIAM IS 2014]

A sequence of measures  $(\mu_n)$  is said to weakly converge to  $\mu$ , if for any bounded continuous function  $\Phi$ ,

$$\int_{\Omega} \Phi(x) \mathrm{d}\mu_n(x) \to \int_{\Omega} \Phi(x) \mathrm{d}\mu(x)$$

Shorthand notation:  $\mu_n \rightharpoonup \mu$ .

### Variable density sampler

A sequence of (random) trajectories  $s_n : X_n \to \Omega$  is said to be a  $\pi$ -Variable Density Sampler if

$$s_{n*}\mu \rightarrow \pi$$
 almost surely

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## Cea vds

![](_page_29_Figure_2.jpeg)

![](_page_29_Figure_3.jpeg)

[Chauffert et al, SIAM IS 2014]

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Cea

## VDS – EXAMPLE (TSP-BASED IN 3D)

![](_page_30_Figure_3.jpeg)

Figure: 3D reconstruction results for r = 8.8 for various sampling strategies. Top row: TSP-based sampling schemes (PSNR=42.1 dB). Bottom row: 2D random drawing and acquisitions along parallel lines [Lustig et al., 2007] (PSNR=40.1 dB).

[Chauffert et al, SIAM IS 2014]

![](_page_31_Picture_0.jpeg)

## **PARAMETERIZATION PROBLEM**

Finding a parameterization in  $S_{MRI}$  corresponding to a curve support is not easy !

• Classical approach, find an admissible parameterization [Hargreaves et al., 2004, Lustig et al., 2008]:

![](_page_31_Figure_4.jpeg)

![](_page_31_Picture_5.jpeg)

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## **PROJECTION OPERATOR**

For an input parameterized curve  $c : [0; T] \rightarrow \Omega$ , define:

$$P_{\mathcal{S}_{\mathsf{MRI}}}(c) = \underset{s \in \mathcal{S}_{\mathsf{MRI}}}{\operatorname{Arg\,min}} \int_{t \in [0;T]} (s(t) - c(t))^2 \mathrm{d}t$$

#### Main properties [Chauffert et al, IEEE TMI 2016]

- Fast resolution using accelerated proximal gradient descent on the dual.
- The sampling time is fixed (equal to T).
- The sampling distribution is well preserved (approximation of Wasserstein distance W<sub>2</sub>).

 $\Rightarrow$  More importantly,  $P_{S_{MRI}}$  is the cornerstone of a global approach, described in part 3.

![](_page_33_Picture_0.jpeg)

## Two key properties for a Variable Density Sampler:

- Sampling distribution
- Fast k-space coverage

## Suboptimal two-step approaches:

• Eg, "Travelling Salesman Problem" sampler

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![](_page_34_Picture_1.jpeg)

## **Part III: Projection on measure sets**

[Chauffert et al, Const Approx 2016; Boyer et al, SIAM IS 2016]

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![](_page_35_Picture_0.jpeg)

## **INTRODUCTION OF A NEW METRIC**

Useful to compare parameterizations and (probability) distributions. Here :  $s : \{1, \ldots, m\} \rightarrow \Omega$  and  $\pi : \Omega \rightarrow \mathbb{R}$  a distribution.

![](_page_35_Picture_3.jpeg)

Related to dithering problem [Teuber et al., 2011].

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## **INTRODUCTION OF A NEW METRIC**

Useful to compare parameterizations and (probability) distributions. Here :  $s : \{1, \ldots, m\} \rightarrow \Omega$  and  $\pi : \Omega \rightarrow \mathbb{R}$  a distribution.

![](_page_36_Picture_3.jpeg)

h: a Gaussian kernel.

### Working with measures

Let  $\mathcal{P}$  denote a set of admissible parameterizations and  $\mathcal{M}(\mathcal{P})$  the set of pushforward measures associated with elements of  $\mathcal{P}$ : Sampling trajectories  $s \in \mathcal{P} \to \Omega$  are seen through  $s_* \mu \in \mathcal{M}(\mathcal{P})$ .

$$\mathcal{M}(\mathcal{P}) = \{ \nu = s_* \mu, \ s \in \mathcal{P} \}.$$

#### *m*-point measures:

Set of sums of *m* Dirac delta functions:  $\mathcal{M}(\Omega^m) = \{ \nu = \frac{1}{m} \sum_{i=1}^m \delta_{p_i}, p_i \in \Omega \}.$ 

Admissible curves for MRI:

$$\mathcal{M}(\mathcal{S}_{\mathsf{MRI}}) = \{ \nu = s_* \mu, s \in \mathcal{S}_{\mathsf{MRI}} \}.$$

We want  $\nu \in \mathcal{M}(\mathcal{P})$  to be "as close as possible to"  $\pi$ , the target distribution.

![](_page_38_Picture_0.jpeg)

## Constructing a metric

Let  $\pi$  denote the *target density*. Let  $\nu$  denote the *pushforward measure*.

Let  $h : \Omega \to \mathbb{R}$  denote a continuous function with a Fourier series that does not vanish. The following mapping:

$$dist(\pi,
u) = \|h\star(\pi-
u)\|_2^2$$

defines a distance (or metric) on  $\mathcal{M}_{\Delta}$ , the space of probability measures on  $\Omega$ .

![](_page_39_Picture_0.jpeg)

Goal: solve numerically, for arbitrary  $\mathcal{M}(\mathcal{P})$ :

$$\inf_{\nu \in \mathcal{M}(\mathcal{P})} dist(\pi, \nu)$$

Theorem [Chauffert et al, Const Approx 2016]

- If  $\mathcal{P} = \Omega^m$ , the sequence of solutions  $\nu_m \rightharpoonup \pi$ .
- If  $\mathcal{P} = \mathcal{S}_{MRI}$ , the sequence of solutions  $\nu_T \rightharpoonup \pi$ .

![](_page_40_Picture_0.jpeg)

### The general construction (similar to finite elements)

• Approximate  $\mathcal{M}(\mathcal{P})$  by a subset  $\mathcal{N}_p \subset \Omega^p$  of *p*-point measures:

$$\mathcal{N}_p = \mathcal{M}(\mathcal{Q}_p) = \left\{ \nu = \frac{1}{p} \sum_{i=1}^p \delta_{q_i}, \quad \text{for } q = (q_i)_{1 \le i \le p} \in \mathcal{Q}_p \right\},$$

where  $\mathcal{Q}_{p}$  is the discretized version of  $\mathcal{P}$ .

• Use a projected gradient descent to obtain an approximate projection  $\nu_p^*$  on  $\mathcal{N}_p$ :

$$\nu_p^* \in \operatorname{Arg\,min}_{\nu \in \mathcal{N}_p} \frac{1}{2} \|h \star (\nu - \pi)\|_2^2,$$

• Reconstruct an approximation  $\nu \in \mathcal{M}(\mathcal{P})$  from  $\nu_p^*$ .

![](_page_41_Picture_0.jpeg)

## NUMERICAL RESOLUTION

## Variational formulation:

$$\min_{\substack{\nu \in \mathcal{N}_p \\ q \in \mathcal{Q}_p}} \frac{1}{2} \|h \star (\nu - \pi)\|_2^2 = \\ \min_{\substack{q \in \mathcal{Q}_p \\ P}} J(q) = \frac{1}{2} \sum_{i=1}^p \sum_{j=1}^p H(q_i - q_j) - \sum_{i=1}^p \int_{\Omega} H(x - q_i) d\pi(x),$$
Repulsion potential
Attraction potential

where *H* is defined by  $\hat{H}(\xi) = |\hat{h}|^2(\xi)$ .

- Repulsion potential: fast k-space coverage
- Attraction potential: right target density  $\pi$
- Generalization of Poisson disk sampling strategy [Bridson, 2007, Vasanawala et al., 2011]

## Projected gradient descents in the non-convex case

Assume that *H* is differentiable with *L*-Lipschitz continuous gradient. Consider the following algorithm:

$$q^{(k+1)} \in P_{\mathcal{Q}_p}\left(q^{(k)} - \tau \nabla J(q^{(k)})\right).$$

The sequence  $(q^{(k)})_k$  converges to a critical point of the functional J.[Attouch et al., 2013].

Remark

In MRI, 
$$P_{\mathcal{Q}_p} = P_{\mathcal{S}_{\mathsf{MRI}}}$$
 !

![](_page_43_Picture_1.jpeg)

## **EXAMPLE : CONTINUOUS LINE DRAWING**

 $\pi = Mona Lisa$ 

![](_page_43_Picture_4.jpeg)

#### Representation of Mona Lisa by $s \in \mathcal{S}_{\mathsf{MRI}}$

![](_page_43_Picture_6.jpeg)

[Chauffert et al, Const Approx 2016]

![](_page_44_Picture_0.jpeg)

![](_page_44_Picture_1.jpeg)

# **Part IV: MRI Simulations Results**

[Boyer et al, SIAM IS 2016]

![](_page_45_Picture_0.jpeg)

# VERY HIGH RESOLUTION IMAGING: SIMULATION SETUP

![](_page_45_Picture_2.jpeg)

#### Parameters:

Image size:  $n = 2048 \times 2048$  (100  $\mu$ m isotropic). m = 0.048n decomposed in:

- 196 radial lines of 1, 024 equispaced samples
- 8 rotated versions of the same spiral made up by 25,000 samples
- 8 curves of 25,000 samples each

![](_page_46_Picture_0.jpeg)

# VERY HIGH RESOLUTION IMAGING: SIMULATION SETUP

![](_page_46_Picture_2.jpeg)

#### MRI hardware constraints:

- $G_{\text{max}} = 40 \text{ mT.m}^{-1} \text{ and } S_{\text{max}} = 150 \text{ mT.m}^{-1} \text{.ms}^{-1}$ .
- For proton imaging,  $\gamma = 42.57 \text{ MHz}.\text{T}^{-1} \implies \alpha = 1,703 \text{ m}^{-1}.\text{ms}^{-1}$  and  $\beta = 6,386 \text{ m}^{-1}.\text{ms}^{-2}.$

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zoom

## VERY HIGH RESOLUTION IMAGING: COMPETING TRAJECTORIES (1/2)

![](_page_47_Picture_2.jpeg)

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Figure : Standard sampling schemes composed of 200,000 samples. (a): i.i.d. drawings. (b): Radial lines; (c): 8 interleaved spirals.

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## VERY HIGH RESOLUTION IMAGING: COMPETING TRAJECTORIES (2/2)

![](_page_48_Figure_2.jpeg)

Figure : Sampling schemes yielded by our algorithm and composed of 200,000 samples. (d): isolated points with repulsion; (e): 8 feasible curves in MRI.

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![](_page_49_Picture_1.jpeg)

## **VERY HIGH RESOLUTION IMAGING CS RESULTS (1/2)**

#### (a) SNR=26.7 dB

![](_page_49_Picture_4.jpeg)

![](_page_49_Picture_5.jpeg)

#### (c) SNR=21.0 dB

![](_page_49_Picture_7.jpeg)

![](_page_49_Picture_8.jpeg)

![](_page_49_Picture_9.jpeg)

![](_page_49_Picture_10.jpeg)

(i.i.d.)

(radial)

(spiral)

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![](_page_50_Picture_1.jpeg)

## **VERY HIGH RESOLUTION IMAGING CS RESULTS (2/2)**

#### (d) SNR=27.0 dB

![](_page_50_Picture_4.jpeg)

#### (e) SNR=23.5 dB

![](_page_50_Picture_6.jpeg)

![](_page_50_Picture_7.jpeg)

#### (*m*-points measure)

![](_page_50_Picture_9.jpeg)

## (admissible curve for MRI)

![](_page_51_Picture_0.jpeg)

## Higher undersampling factors achieved at higher resolution

- Up to 20-fold acceleration at 100  $\mu m$  in-plane

## Better image quality achieved using projection on measure sets

- Best results given by the projection on *m*-points measures
- Projection on admissible curves for MRI outperforms radial and spiral sampling schemes by 2-3 dB

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![](_page_52_Picture_1.jpeg)

## Part V: Retrospective & Prospective SPARKLING at 7T

**SPARKLING**: Segmented Projection Algorithm for Random K-space sampLING

[Lazarus et al, submitted to ISMRM'17 & to IEEE TMI] [Lazarus et al, in prep. to MRM]

![](_page_53_Picture_0.jpeg)

 SPARKLING <u>outperforms classical</u> sampling schemes (eg, radial, spiral) in simulations but in the real life?

![](_page_53_Figure_3.jpeg)

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## **ADAPT TRAJECTORIES TO MR PHYSICS**

1) Shorter observation time & optimized T2\* contrast

- Shorter  $T_{obs} \approx 35 \text{ ms}$
- Echo time adapted to contrast: TE ≈ 30 ms (directed trajectories)

![](_page_54_Figure_5.jpeg)

- 64 segments lasting 33.3 ms and collecting 1024 ADC samples each.
- Each segment (one is in blue) passes trough the kspace center at echo time TE=30 ms.
- Gmax = 40 mT/m
- Smax = 200 T/m/s

![](_page_55_Picture_1.jpeg)

#### 2) Check for gradient errors

Goal: Estimate gradient errors on novel trajectories.

Methods: LPM (Local Phase Measurement) to measure actual gradients values

![](_page_55_Figure_5.jpeg)

N = 256 R = 4

#### **2)** Check for gradient errors

**Results:** The measured k-space locations are **very close** to the prescribed sampling scheme. Largest errors are observed in regions of high curvatures.

![](_page_56_Figure_4.jpeg)

![](_page_57_Picture_0.jpeg)

## **EXPERIMENTS**

• Ex-vivo brain baboon in fluorinert solution

#### Sequence parameters

- 7T Siemens healthineers scanner
- In vivo Corp Birdcage 1Tx/1Rx coil
- T2\* weigthing
- Slice thickness: 3 mm
- TR = 60 ms
- TE = 30 ms
- $-\alpha = 25^{\circ}$
- Axial slice
- Signal averaging to increase input SNR

![](_page_57_Picture_13.jpeg)

![](_page_57_Picture_14.jpeg)

Full Cartesian acquisition at N=512 and N=1024

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## **RESOLUTION DEPENDENCE**

![](_page_58_Figure_2.jpeg)

# 

# **RESOLUTION DEPENDENCE: RETROSPECTIVE EXPERIMENTS**

N = 512 Input SNR = 78

![](_page_59_Picture_3.jpeg)

Reference

![](_page_59_Picture_5.jpeg)

SSIM = 0.83

SSIM = 0.76

![](_page_60_Figure_0.jpeg)

## **RETROSPECTIVE CS RESULTS – N = 1024**

![](_page_60_Figure_2.jpeg)

![](_page_61_Picture_0.jpeg)

### **Full Cartesian sampling**

#### 8-fold acceleration

![](_page_61_Picture_4.jpeg)

![](_page_62_Picture_0.jpeg)

## • Retrospective CS results:

- SPARKLING trajectories outperform Golden Angle radial ones on real data at 7 Tesla
- Are consistent with synthetic simulations: **16-fold** acceleration
- Confirm the dependence of the acceleration factor on the pixel size (or image resolution defined by k<sub>max</sub>)

#### • Prospective CS results

- Our CS-GRE T2\*-weighted sequence works!
- Lower acceleration achieved so far (8-fold) as compared to retrospective CS
- Gradient imperfections and  $B_0$  inhomogeneities must be accounted for.

## ACKNOWLEDGEMENTS

#### Nicolas Chauffert

CEA/NeuroSpin

![](_page_63_Picture_4.jpeg)

Carole Lazarus CEA/NeuroSpin

![](_page_63_Picture_6.jpeg)

Alexandre Viggnaud CEA/NeuroSpin

![](_page_63_Picture_8.jpeg)

#### Pierre Weiss CNRS/ITAV

![](_page_63_Picture_10.jpeg)

Claire Boyer IMT, U. Toulouse

![](_page_63_Picture_12.jpeg)

Jonas Kahn CNRS/IMT

![](_page_63_Picture_14.jpeg)

![](_page_64_Picture_1.jpeg)

# Thanks for your attention! Any questions?