## Head motion in diffusion MRI

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#### Diffusion contrast

- Basic principle of diffusion MRI: Microscopic motion of water molecules causes attenuation of MR image intensity
- {DW image intensity} = {Baseline image intensity}
  × {Attenuation factor}
- All diffusion MRI models are based on this, *e.g.*:



• But macroscopic head motion also acts this way!

- Head motion during a dMRI scan can lead to:
  - Misalignment between consecutive DWI volumes in the series
  - Attenuation in the intensities of a single DWI volume/slice, if the motion occurred during the diffusion-encoding gradient pulse
  - The former can be corrected with rigid registration, *the latter can't*



- Conventional EPI sequences for dMRI ignore the problem
  - If motion in several directions  $\Rightarrow$  underestimation of anisotropy
  - False positives in group studies where one group moves more
  - Effects more severe when higher *b*-values, more directions acquired

## Motion in a dMRI group study

50 children with autism
 spectrum disorder (ASD) and 62
 typically developing children
 (TD), ages 5-12

- 165 total scans (some retest)
- DWI: 3T, 2mm isotropic, 30 directions, b=700 s/mm<sup>2</sup>
- Outlier data sets excluded
- Pathways reconstructed with TRACULA



Yendiki et al., 2013

Data courtesy of Dr. Nancy Kanwisher and Ellison autism study

#### Between-volume motion measures

Yendiki et al., 2013

- From eddy-current correction: affine registration of each volume to the first baseline volume
- Find rotation and translation
- Transform to relative rotation and translation from each volume to the previous volume



• Measure 1: Average volume-by-volume translation

 $- \Sigma_{k} \operatorname{sqrt}(x_{k}^{2} + y_{k}^{2} + z_{k}^{2}) / N_{vol}$ 

- Measure 2: Average volume-by-volume rotation
  - $\Sigma_k(|\theta_k| + |\phi_k| + |\psi_k|) / N_{vol}$

#### Within-volume motion measures

Yendiki *et al.,* 2013

- Score of DWI intensity drop-out [Benner *et al.*, MRM 2011]:
  - For each volume, define an intensity threshold: *I<sub>min</sub>* = *T* · exp(-*b*·*D*)
    *T* the intensity threshold for brain voxels at *b*=0 (default: *T* = 100)
    *b* the b-value of the current volume

*D* a nominal value of diffusivity in the brain (default: D = 0.001)

- Motion score:  $S = 2 r / (0.7 \cdot r_1)$ 
  - **r** the number of voxels with intensities  $I > I_{min}$  in this volume
  - **r**<sub>1</sub> the number of voxels with intensities  $I > I_{min}$  in the first volume with the same b-value
- S > 1 : Excessive drop-out
- Measure 3: Percentage of slices with signal drop-out
  - Number of slices with S > 1 over the total slices in the data set
- Measure 4: Signal drop-out severity
  - Average score S for slices with S > 1

# Motion measures histograms

Yendiki *et al.,* 2013

- Qualitative visual inspection to find scans with excessive motion
- Of the 165 scans, 17 were removed after inspection



## Motion measures by group

Yendiki *et al.,* 2013

• Significant differences in median measures between groups



For ASD children, rotational motion is correlated with ADOS score (*ρ*=0.31, *p*=0.04)

#### Yendiki *et al.,* 2013

- 50,000 random combinations of scans from 30 ASD and 30 TD children, agematched
- Compute difference in motion measures between ASD and TD
- Count the tracts that have significantly different FA between ASD and TD





Yendiki *et al.,* 2013

#### Example trial:

Lower motion difference, one FA finding



#### Example trial:

Higher motion difference, six FA findings



#### Yendiki *et al.,* 2013

#### Differences in anisotropy and diffusivity measures between groups

500 groups with lowest differences in head motion:

- Translation 0.04 ±0.03 mm
- Rotation: 0.05  $\pm$  0.01  $^{\circ}$
- Portion of slices with drop-out:  $0.04 \pm 0.02 \%$
- Drop-out score:  $0.02 \pm 0.02$

- Translation:  $0.36 \pm 0.03 \text{ mm}$
- Rotation: 0.28  $\pm$  0.01  $^{\circ}$
- Portion of slices with drop-out:  $0.11 \pm 0.02 \%$
- Drop-out score:  $0.08 \pm 0.02$



#### Yendiki *et al.,* 2013

#### Frequency of significant (p < 0.05) group differences

#### 500 groups with lowest differences in head motion:

- Translation 0.04  $\pm$  0.03 mm
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#### Motion as a nuisance regressor

Yendiki *et al.,* 2013

• Total motion index (TMI) of the *i*-th scan:

 $\text{TMI}_{i} = \sum_{j=1,...,4} (x_{ij} - M_{j}) / (Q_{j} - q_{j})$ 

where:  $x_{ij}$  each of the 4 motion measures for this scan  $M_j$  its median value over all scans  $Q_j$  its upper quartile over all scans  $q_j$  its lower quartile over all scans

• Compute the TMI of each scan, include it as a regressor in the group analysis

#### Yendiki *et al.,* 2013

#### Frequency of significant (p < 0.05) group differences, TMI regressed

#### 500 groups with lowest differences in head motion:

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# TD vs. TD

#### Yendiki *et al.,* 2013

#### Differences in anisotropy and diffusivity measures between groups

500 groups with lowest differences in head motion:

- Translation  $0.04 \pm 0.03 \text{ mm}$
- Rotation: 0.0003  $\pm$  0.0002  $^{\circ}$
- Portion of slices with drop-out:  $0.01 \pm 0.01 \%$
- Drop-out score:  $0.02 \pm 0.01$

- Translation:  $0.28 \pm 0.05$  mm
- Rotation: 0.20  $\pm$  0.01  $^{\circ}$
- Portion of slices with drop-out:  $0.07 \pm 0.01 \%$
- Drop-out score:  $0.07 \pm 0.02$



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#### Test vs. retest

Yendiki *et al.,* 2013

Lower-motion vs. highermotion scans of *the same* 25 TD children



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#### Voxel-based analysis

Yendiki *et al.,* 2013

• Test each voxel for association of FA with head motion with TBSS [Smith '06] on all scans



#### Replication in adult data

• Test each voxel for association of FA with head motion with TBSS on data from a group of middle aged and older adults



Salat, Chapter 12 in: *Diffusion MRI*, 2nd edition (in press)

#### Motion compensation strategies

- Retrospective:
  - Registration-based [Andersson '02, Rohde '04]
    - Does not correct for intensity drop-out, less robust at high b-values
  - Outlier removal [Chang '05, Zwiers '10]
    - Must have redundancy in data, remove comparably from every group
  - Nuisance regressors
  - Motion matching between groups
- Prospective:
  - Motion-compensated sequences
  - Accelerated sequences

# Data acquisition solutions: Motion-compensated diffusion MRI



# Motion-compensated dMRI

Work by Himanshu Bhat

- Optical tracking systems [Aksoy '11]
- Selective reacquisition [Benner '11]
- Free induction decay navigators [Kober '12]
- Volumetric navigators:
  - Low-res, non-DW EPIs acquired between TRs [Alhamud '12] or between slices throughout each TR [Bhat '12]
  - No external hardware needed
  - Performance independent of b-value
  - Small contribution to TR



#### dMRI with volumetric navigators

Work by Himanshu Bhat



## dMRI with volumetric navigators

Work by Himanshu Bhat



#### dMRI with volumetric navigators

Work by Himanshu Bhat

- Resolution
  2mm isotropic
- TR = 7.3 s
- TE = 69 ms
- 100 directions
- *b*=1000 s/mm<sup>2</sup>



# Data acquisition solutions: Accelerated diffusion MRI



## Simultaneous multi-slice EPI

- Motivation: Increase efficiency of EPI for dMRI and fMRI
  - Whole brain acquisition with thin slices leads to long TR
  - TR >> T1  $\Rightarrow$  inefficient SNR/time
  - Long acquisition time makes EPI with high spatial resolution (or, for dMRI, high angular resolution) impractical
- Conventional parallel imaging: Undersample data in *k*-space
  - For *R*-fold speedup, SNR drops by  $g\sqrt{R}$
  - Shorter read-out train but efficiency gain smaller for EPI
- Simultaneous multi-slice (SMS): Excite and acquire multiple slices in each readout period
  - For *R*-fold speedup, SNR drops only by g
  - Reduces TR significantly



# Teasing apart the slices

- Parallel imaging with simultaneous acquisition of multiple slices:
  - High *g*-factor when slices are close to each other [Larkman '02]
- CAIPIRINHA:
  - Fix this for multi-slice parallel FLASH with FOV/2 shifts b/w slices by phase cycling of excitation pulses [Breuer '05]
  - Not relevant to EPI
- Extension to EPI:
  - Use blips in slice-select direction to shift slices to get benefits of CAIPIRNHA [Nunes '09]
  - Constant slice-select blips with every phase-encode blip cause through-plane dephasing
    - $\Rightarrow$  "tilt" voxels in phase-encode direction
- Blurring solved by blipped-CAIPI:
  - Back-and-forth jumps in slice-select blips to unwind dephasing of previous blip [Setsompop '12]







# Simultaneous multi-slice dMRI

- Multi-slice SE-EPI for diffusion imaging with 3-fold speed-up
- Resolution 2mm
  isotropic
- 32-channel head coil
- Inter-slice gap 40mm
- FOV 208 x 208 x 120mm
- TR = 3s
- TE = 96 ms
- Partial Fourier <sup>3</sup>/<sub>4</sub>
- Matrix 104x78x20
- Slice GRAPPA



## Simultaneous multi-slice dMRI

Work by Kawin Setsompop

- DTI (64 directions,  $b=1000 \text{ s/mm}^2$ , 2 mm isotropic)
  - Conventional acquisition: 10 min
  - 3-fold SMS acquisition: 3 min



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#### Simultaneous multi-slice dMRI

Work by Kawin Setsompop

- DSI (257 directions,  $b_{\text{max}}$ =7000 s/mm<sup>2</sup>, 2.5 mm isotropic)
  - Conventional acquisition: 41 min
  - 3-fold SMS acquisition: 14 min

# tx acquisition 3x acquisition

# Simultaneous multi-slice fMRI

- 7T fMRI, 1mm isotropic, 120 slices (whole head), R=2 in-plane GRAPPA, 32 channel coil
- TR=2.88s with 3x simultaneous multi-slice acquisition



#### Conclusions

- Differences in head motion between groups can induce spurious group differences in diffusivity and anisotropy
- General trend: Head motion  $\uparrow \Rightarrow RD\uparrow$ ,  $AD\downarrow$ , MD -,  $FA\downarrow$
- This is *after* registration-based motion correction
- Match motion between groups and/or use a motion score as a nuisance regressor
- Note that all this will address *false positives*, but not *false negatives* due to head motion in the data
- Methods for tackling the problem during data acquisition are important

