## DECLARATION OF CONFLICT OF INTEREST OR RELATIONSHIP

Speaker Name: Terry Oakes
I have no conflicts of interest to disclose with regard to the subject matter of this presentation.

## Proronto!

# Artifacts in FMRI 

Terry Oakes<br>University of Wisconsin-Madison<br>Waisman Center Brain Imaging Lab

WISCONSIN

What I can tell you about:

- PET imaging
- Radionuclide production
- PET data analysis
- fMRI data analysis
- Coregistration
- Morphometry measures


Not so much:
MRI physics, pulse sequences, data acquisition

## What is an Artifact?

## To neuroscientists, this brain looks funny.

To Wisconsin Badger fans, the brain obscures the logo.

An artifact is anything that prevents you from seeing what you want to see.

## What is the Signal?

Timeseries - filtered_func_data (face/voice)



## What is the goal?



## Big-picture goal



## Artifact Sources



## A Scientific Experiment

- Before
- measure twice, cut once
- During

Q look at your data. with your eyes.


- After

0 pre-processing: modeling and removing variance
D data analysis: modeling and discovering signal

## The single biggest artifact source



## Being Clever Ahead of TIme

- Before
- Learn a package

O provides an orderly introduction to analysis.

- Software programming skills
- File format conversion
- Experimental paradigm

Q event-related paradigms can isolate stimulus, HR

- Acquisition sequence

DCollect enough data
O multispectral anatomicals (T1, T2, PD, ...)
Ofield map
O physiological (cardiac, respiratory, skin conductance, eye tracking)
Destimator for HRF (e.g. motor response)

## Post-hoc Corrections

- Some folks call it image processing...
- spatial registration
- field map correction
- slice time correction

D motion correction

- smoothing- spatial, temporal
- These corrections are ignorant of the experimental model.


## EPI Dropout (susceptibility) artifact


phase dispersion => signal loss

## EPI susceptibility artifact

clever ahead of time: optimize acquisition parameters



## EPI registration - tread carefully



## EPI registration: indirect approach



## Typical acquisition sequence

- Goal: spatial alignment
- Hurdle: movement ~ time



## Off to a bad start




Remove first 3-5 frames:
D delete from series or

- mark via GLM software
or
- assign to a unique condition


## Field Map Correction

- The magnetic field is not uniform, leading to misplaced signal, since recording/reconstruction assumes a uniform gradient.
- $3 \mathrm{~T}=>$ worse
- Nonuniformities are object-dependent.
- Largest source of $B_{0}$ inhomogeneity: air-tissue interfaces $=>$ susceptibility differences
- Different acquisition sequences may have different affect.

O EPI data (Gradient Echo) most affected (fMRI, DTI)
O Spin Echo affected little
O Luckily, this can be modeled and corrected...
O But, a seperate measurement is required.
O 2 scans with different TEs (e.g. 8, 11 msec ), minimal distortion.

## B-Field Distortion

## The Cause: EPI Phase Error Accumulation



- For a linear field offset, phase error accumulates linearly.
- Cummulative phase errors cause a shift in position


## Distortion (Fieldmap) Correction



- Strategy: Collect two gradient echo images at short but different echo times.
- Unwrap the phase.

Q Phase-difference proportional to the pixel-offset along the phase-encode axis.

- Resample the EPI image to correct for the offset.

$\mathrm{TE}=7 \mathrm{~ms}$

$\mathrm{TE}=10 \mathrm{~ms}$

Peter Jezzard and Stuart Clare, "Sources of distortion in functional MRI data", Human Brain Mapping, 8:80-85, 1995.

## Field Map Correction



## Field Map Correction

Magnitude of correction depends on the magnitude of the susceptibility artifact, which depends on the size of the sinuses.

OLarger in men than women.

- Larger in adults than children
- Although the magnitude of the correction is often small, it can reduce confounds between gender, age and susceptibility.

OIt cannot recover missing data from the dropout region.


## Inhomogeneity Correction

- Increase gradient magnetic field strength.

Decrease echo time.

- Smaller pixels (better resolution).
- Phase encoding.
- Postprocessing.



## Slice-time Correction



Make all slices appear to have been acquired at the same time.

A "reference" HRF should look the same in all slices.


More important for longer TR. Usually best prior to motion correction.

## Motion Artifacts

- Sources
- Subject motion
- Peripheral movement (changes B-field)
- Respiration, cardiac
- Magnitude / Importance

Q a significant fraction of the fMRI signal!

- Model
- framewise 3D volumetric

O rigid body (translation, rotation)
Oignores intraframe motion
Q assumes only small movement

- Correlation between motion, stimulus

D correcting motion can attenuate activation signal
D motion estimates can be incorporated into GLM

## Motion Correction: Magnitude



## Motion Correction



## Image realignment

$1 /$ scratch/TMRI_data/data/MRI_vis_stim_0000.img
$2 /$ scratch/MRI_data/data/MRI_vis_stim_0001.img
3 /scratch/MMRI_data/data/MRI__vis_stim_0002.img
4 /scratch/MRI_data/data/MRI_vis_stim_0003.img
5 /scratch/MRRI_data/data/MRI_vis_stim_0004.img
6 /scratch/MRRI_data/data/MRI_vis_stim_0005.img
/scratch/MMRI_data/data/MRI_vis_stim_0006.img
3 /scratch/MRRI_data/data/MMRI_vis_stim_0007.img
9 /scratch/MMRI_data/data/MRI_ vis_stim_0008.img
10 /scratch/TMRI_data/data/MRI_vis_stim_0009.img
$10 /$ scratch/MRI_data/data/MRI_vis_stim_0009.img
$1 /$ scratch/TMRI_data/data/TMRI_Vis_stim_0010.img
12 /scratch/TMRI_data/data/MRRI_vis_stim_0011.img
...........etc


## Motion Correction: Subject Motion


t=later

between-plane: messy



## Combining scan runs



## Motion Artifact: Typical pattern

Single subject data
effect size


## Motion Correlated with Stimulus



## Motion Parameters:

## How can we use this information for Good?

-Apply motion correction: reslice each 3D volume
${ }^{\prime \prime}$ standard" aproach
-loss of sensitivity if motion correlated with activation
OUse parameters as covariates in GLM
. may increase sensitivity

- more flexible data analysis
. small loss of degrees of freedom in GLM


## Motion Parameters as GLM Covariates

## -Block design:

. if motion correlated with stimulus, "standard" reslice best -Event-related design:
. motion parameters in GLM usually work well.


## Motion Correction at Work



## Smoothing - spatial

OIncrease S:N

- Objects similar in size to smoothing kernal emphasized
- matched filter theorem
- Manage imperfect registration
©Fulfill "Gaussian random field" assumption



## Smoothing

## 



## Smoothing - when?

- EPI time series

O Spatial smoothing removes small (uninteresting) clusters
O Mild temporal smoothing beneficial:
a trend removal
4igh-pass filter
Q most versatile if left until GLM stage

- Parameter estimate (cluster) maps

O more versatile for analysis
Ochanges fitted results

## Group-wise artifacts

O Multi-subject or 2cnd-level analysis
O subject-to-subject variance dominates
O most analysis assume:
D similar within-subject variance

- similar data acquisition, analysis



## Software Recommendations

- fMRI - specific
- AFNI, FSL, SPM have largest market share, similar results
- Two- step procedure for selecting software:
- Find the smartest person in your lab.
- Use what they use.
- Scripting
b bash
- fast, easy, universal
- idiosyncratic, hard to distribute
- Python

Q versatile, extensible, good for distribution
O mostly universal, moderate learning curve
D Matlab, IDL
Q sed \& awk? You don't need my advice.

- Other handy tools:
- R

D fmristat

- NiPy
http:/ / www.r-project.org/
http:// www.math.mcgill.ca/ keith/fmristat/
http:// neuroimaging.scipy.org/

