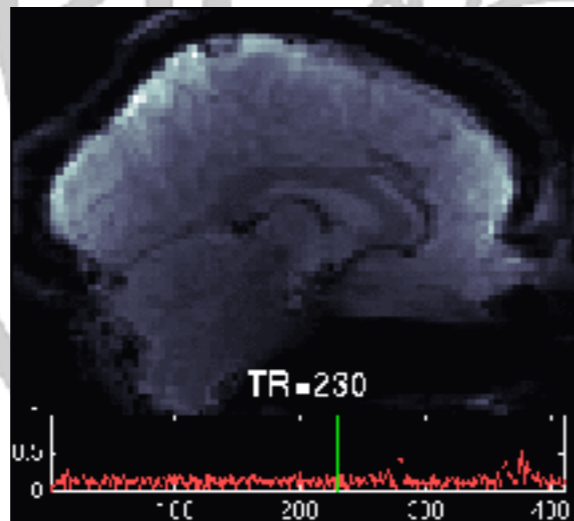


# Assessment of “denoising” (motion artifacts removal) on resting state fMRI data



Method Club

2015-10-15

Seung-Goo (“SG”) KIM

# Questions to discuss today

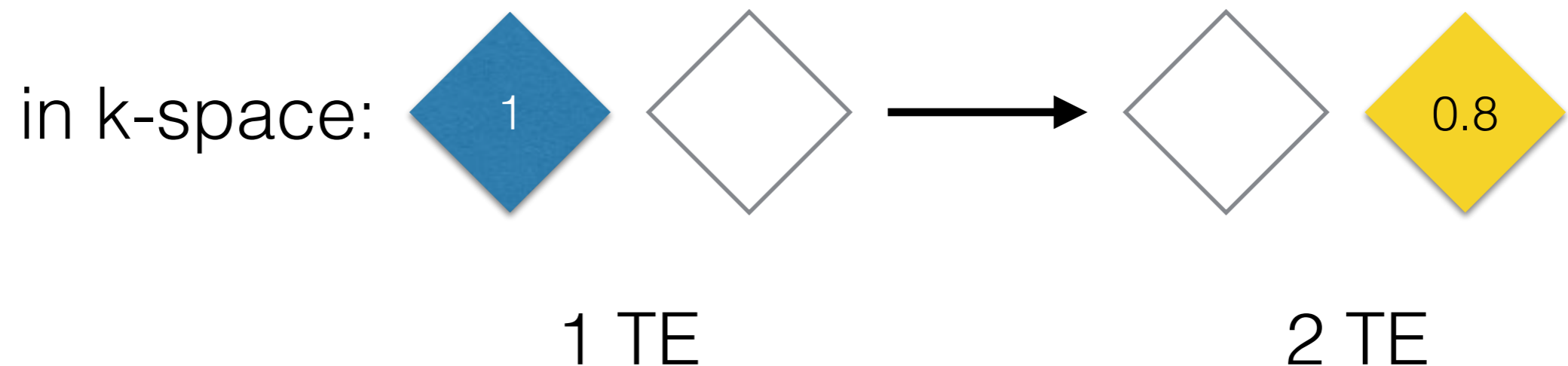
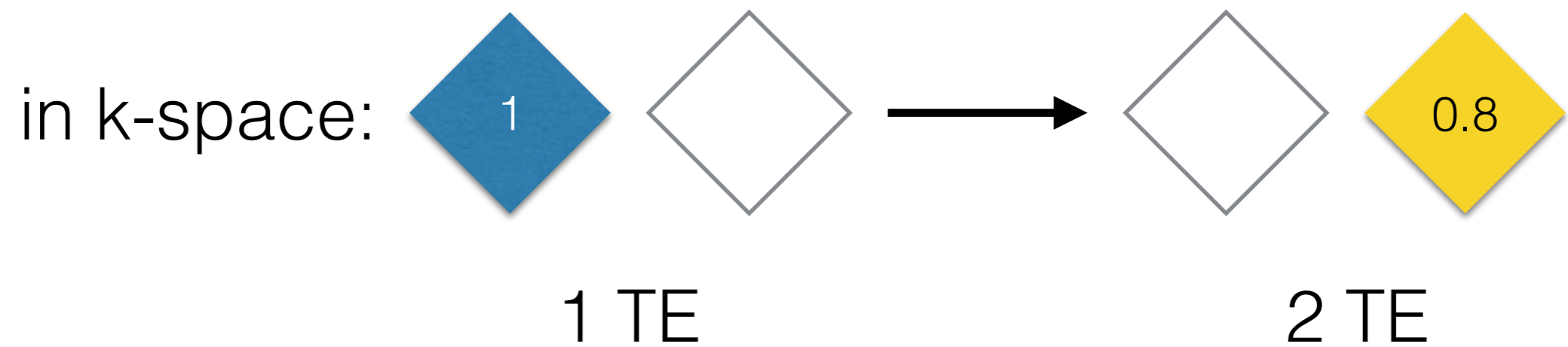
1. Why do you want to 'modify' your data?
2. If you do it, how can you tell it's improved or worsened?
3. Which parameters work better than others?

\*DISCLAIMER: This is not to defend/recommend a certain toolbox (e.g. CONN) but to openly discuss about the assessment of denoising process for rs-fMRI!

# Question #1

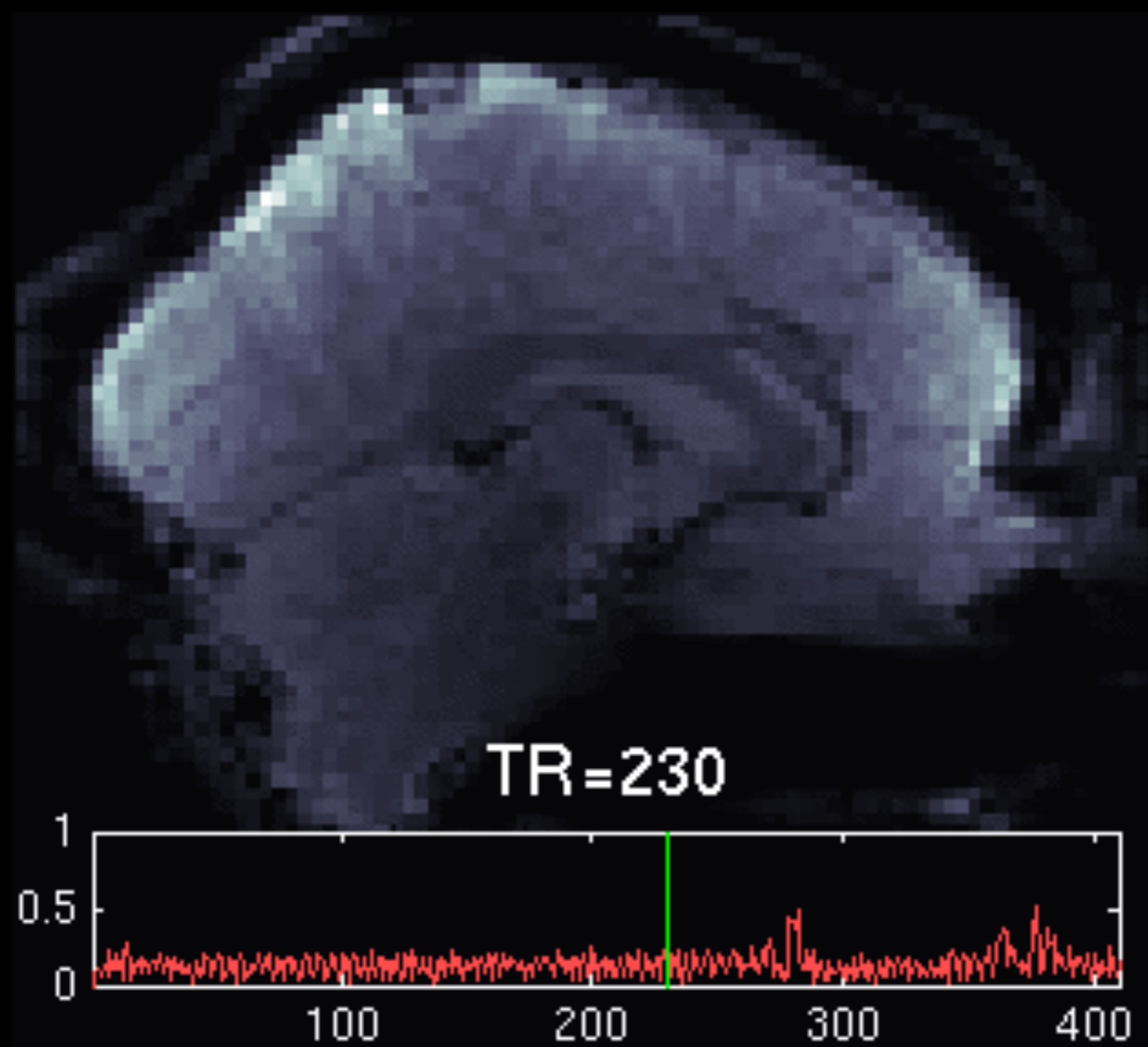
- Why do we have to “modify” our data?
- Because head motion does not only change the position, but also change IMAGE INTENSITY too!
- Head motion particularly creates ***synchronized signal change***, which results in heightened correlation over nearby (and also distant) voxels.

# Why signal changes?

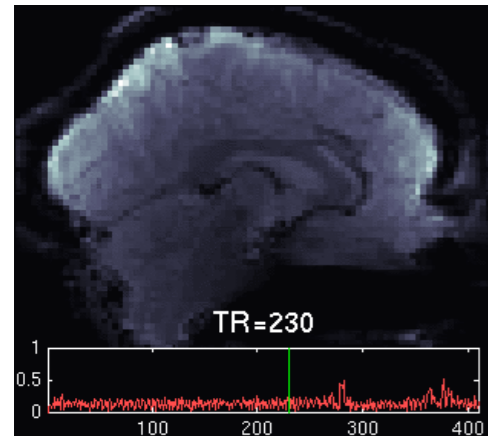


# Example

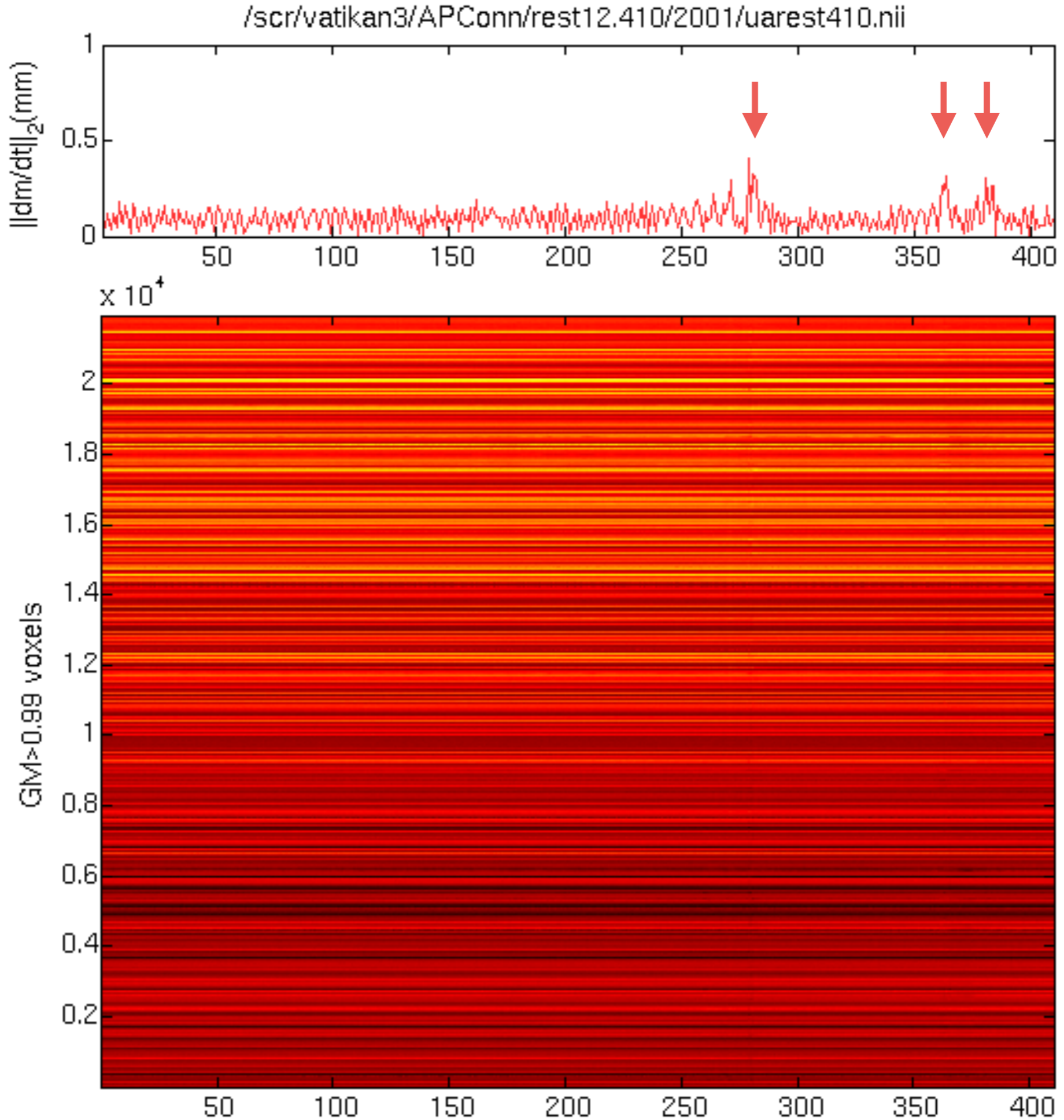
- MAGNETOM Prisma at 3-T
  - 4x multiband EPI sequence of 64 axial slices with 88 x 88 image matrix (“LEMON” sequence)
  - TR/TE= 1400/30 msec; FA= 69 degrees
  - 420 volumes (9.8 min)
  - Voxel size= 2.295 x 2.295 x 2.300 mm<sup>3</sup>
- Subject: a healthy male musician (German)



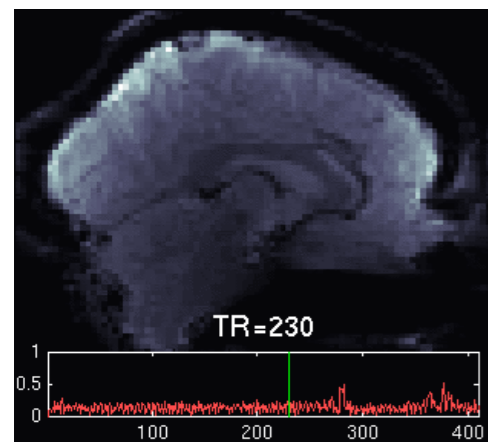
even after realignment



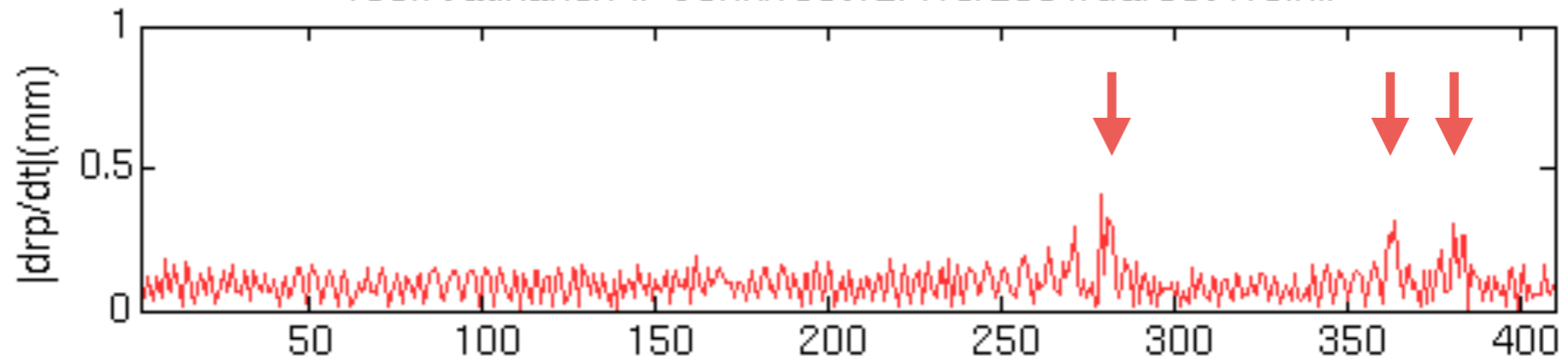
# Image Intensity



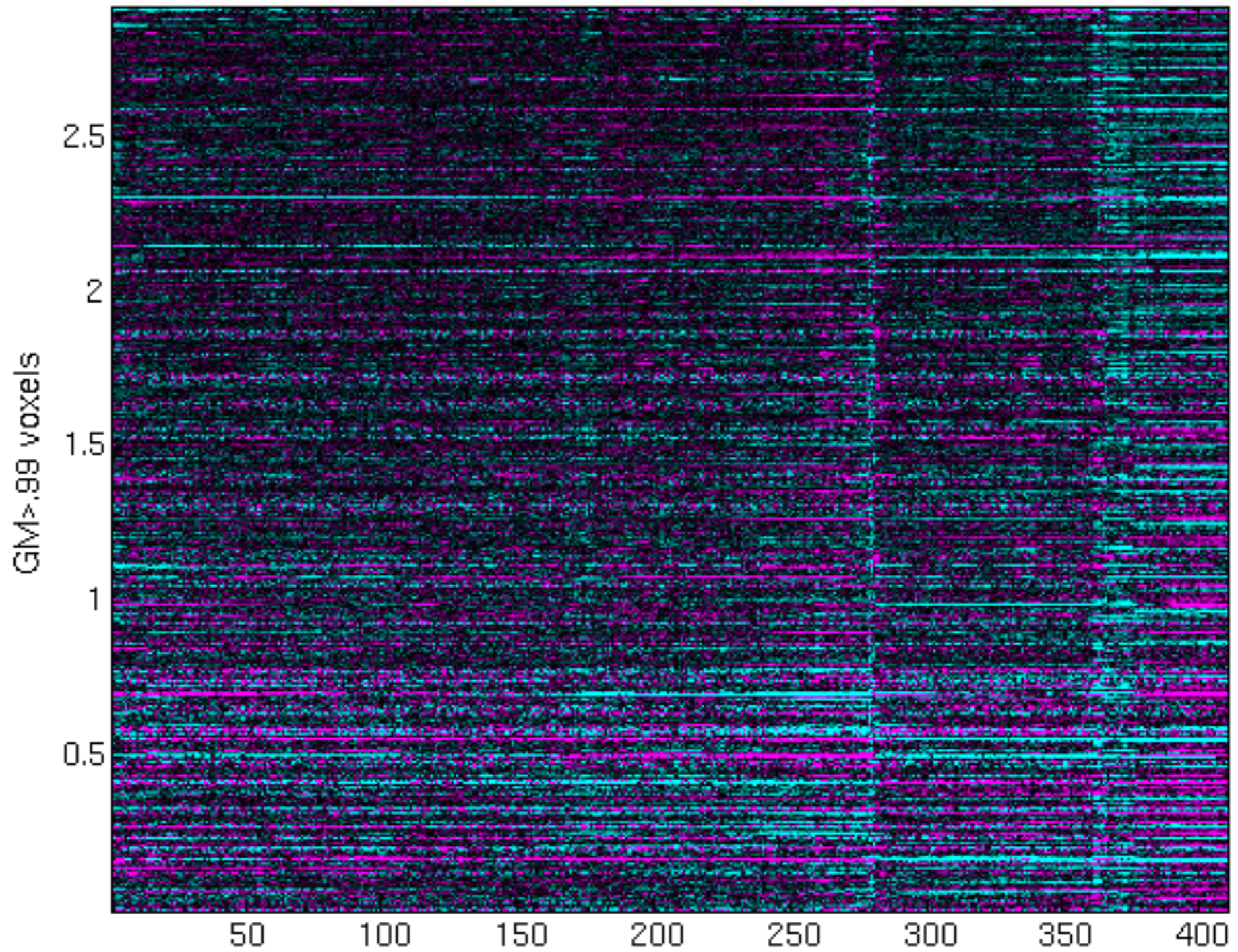




/scr/vatikan3/APConn/rest12.410/2001/uarest410.nii



$\times 10^4$



Signal change from mean:

$$\frac{x(t) - \bar{x}}{\bar{x}}$$

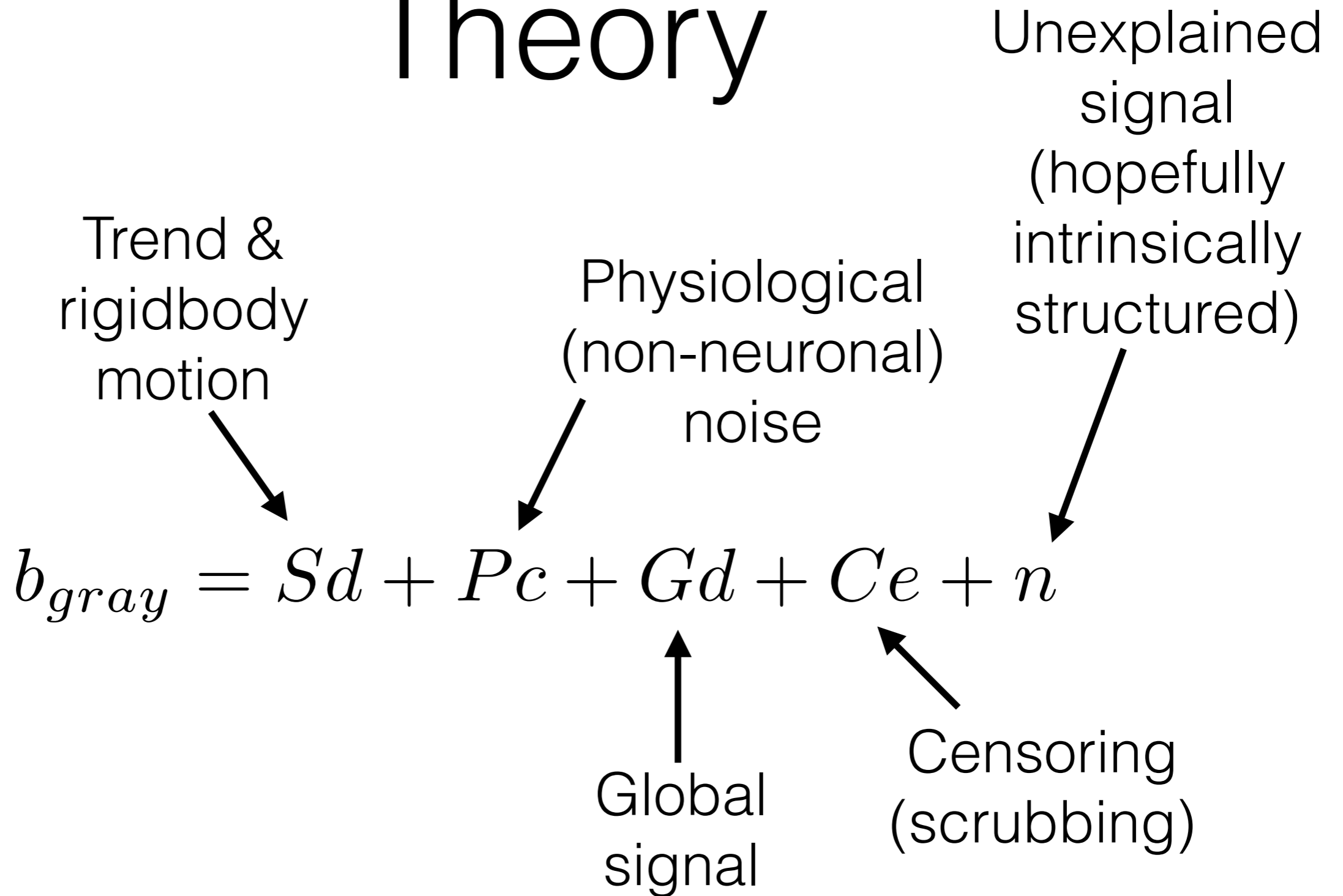
Change from mean(%)



# Question #2

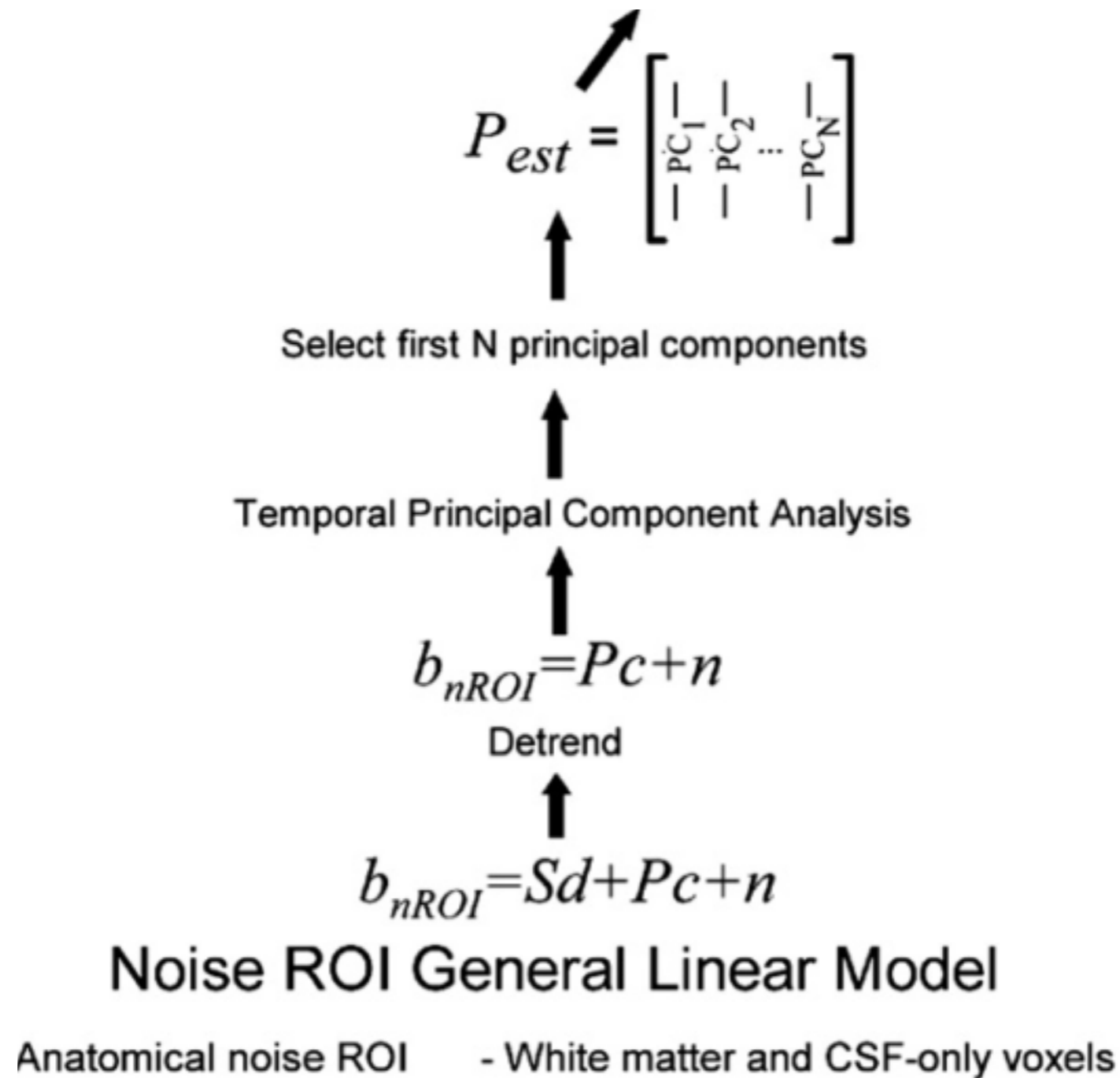
- How can we tell whether some signal is induced by head motion or neural activities?
- We (I) assume:
  - head movements affects extensively (global signal)
  - WM/CSF voxels doesn't show neuronal hemodynamics but motion-induced signal change (CompCor)
  - correlation between random GM voxels would be close to Gaussian, at least under the null model (K-S test)

# Theory



(modified from Behzadi et al., 2007, NI.)

# Physiological noise?



Anatomical noise ROI - White matter and CSF-only voxels

# “Denoised” timeseries

$$b_{gray} = Sd + Pc + Gd + Ce + n$$

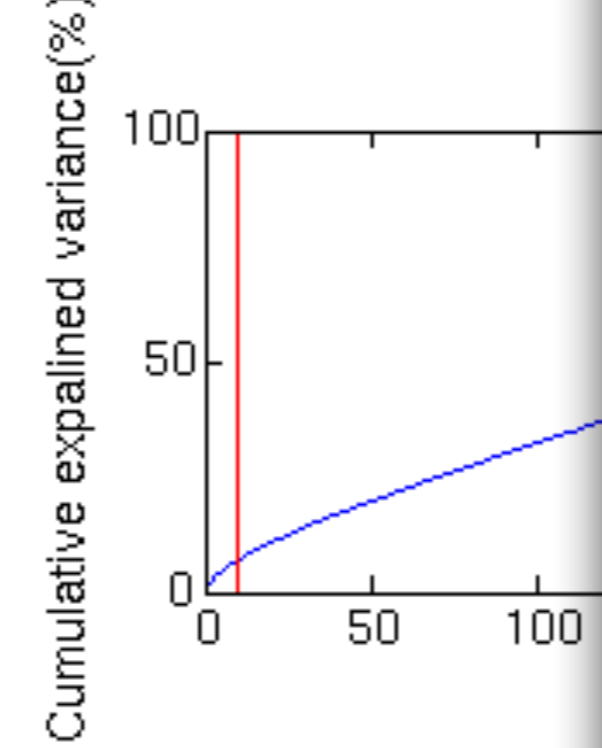
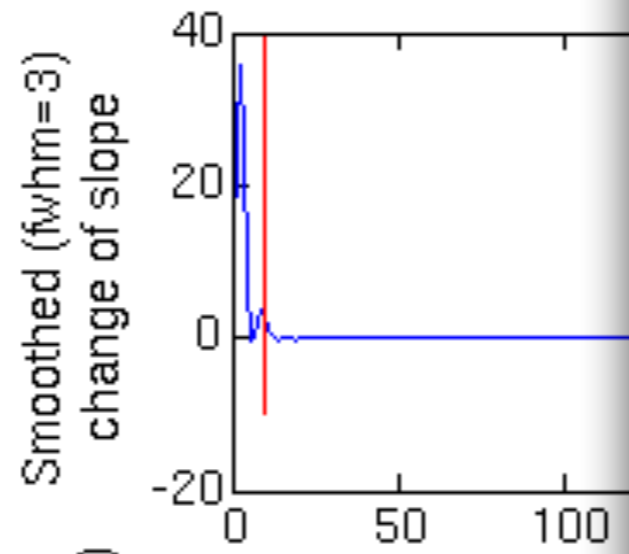
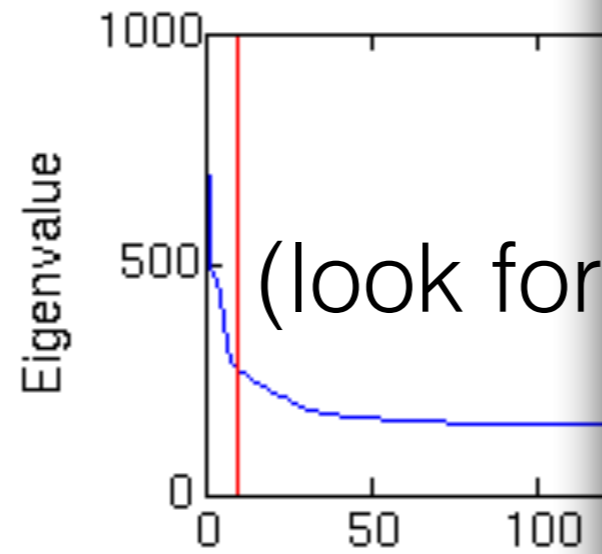
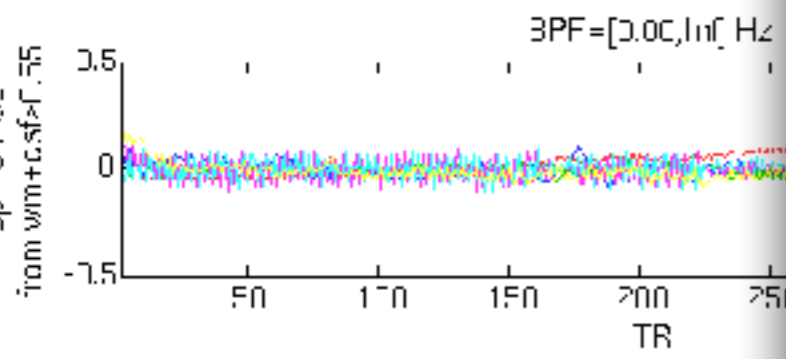
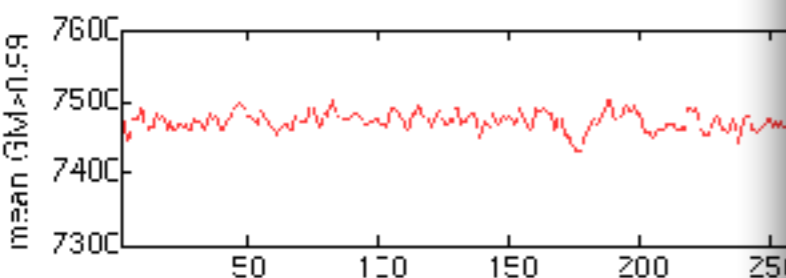
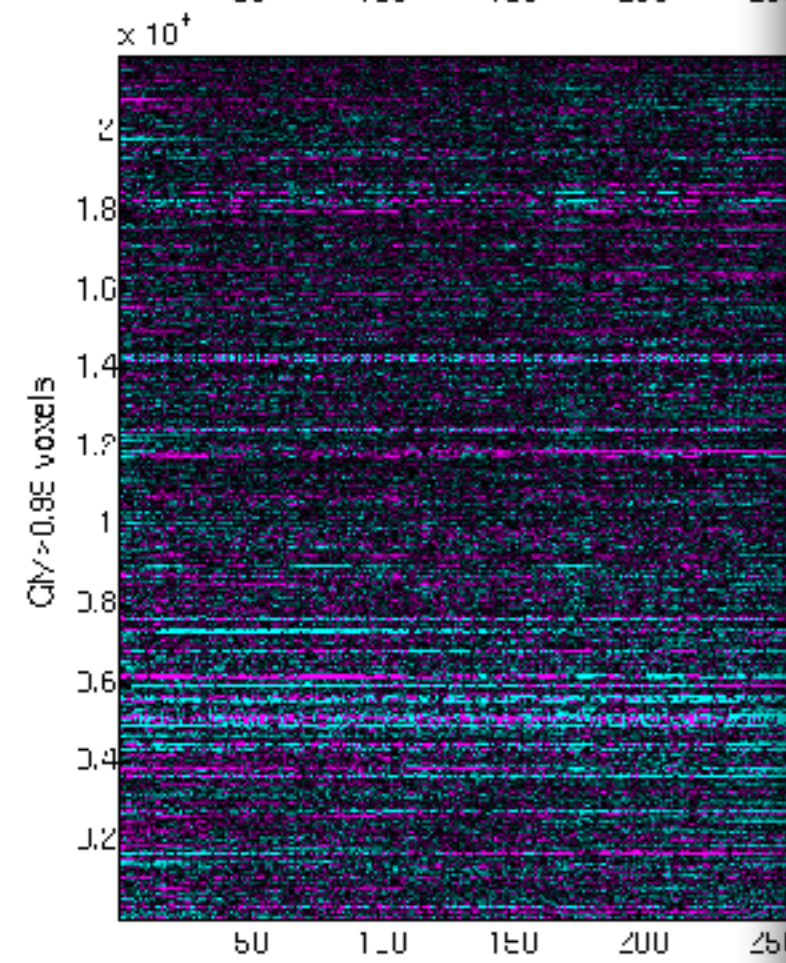
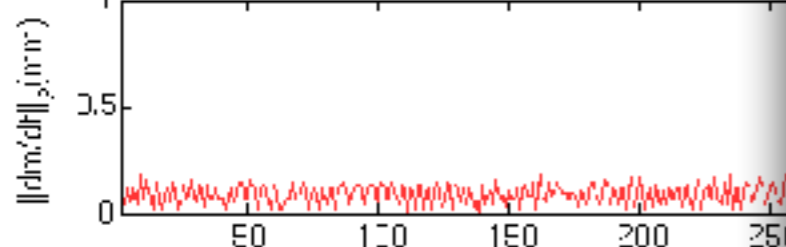
$$\hat{b}_{gray} = S\hat{d} + P\hat{c} + G\hat{d} + C\hat{e}$$

$$\text{residual} = b_{gray} - \hat{b}_{gray}$$

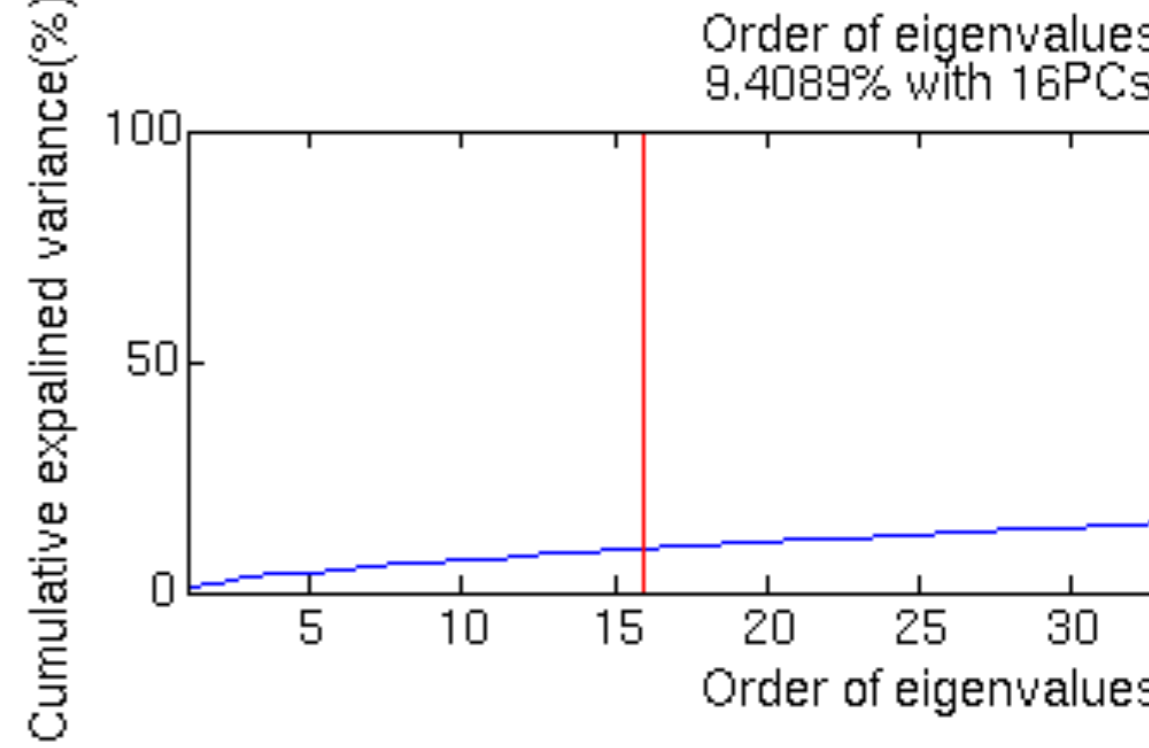
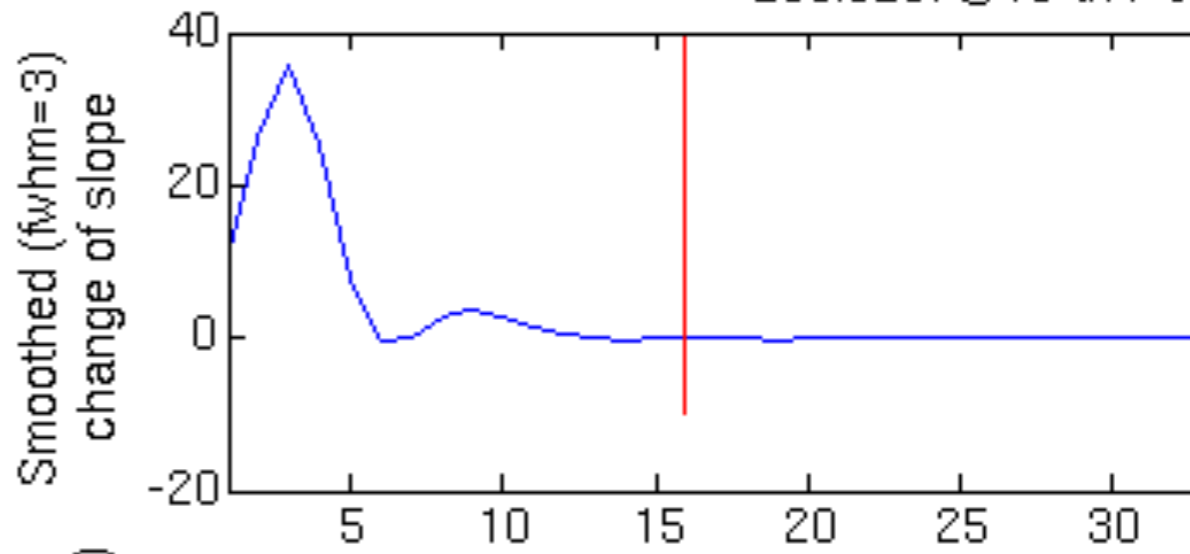
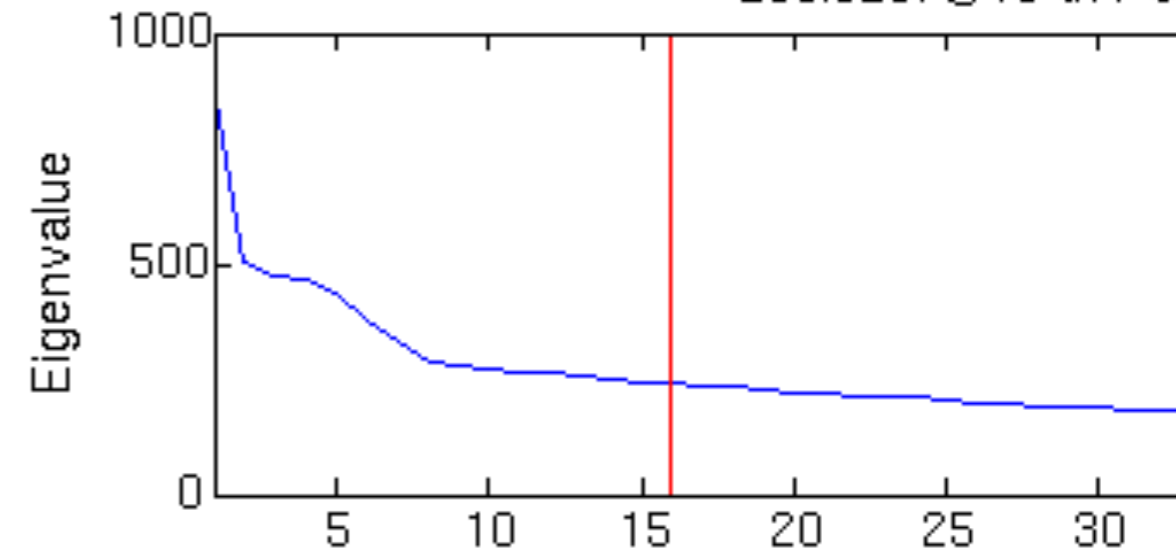
...really?

Visual inspection

ispr/vetikan/VAPConn/rest12.41 V20



239.0207@16-th PC



(look for

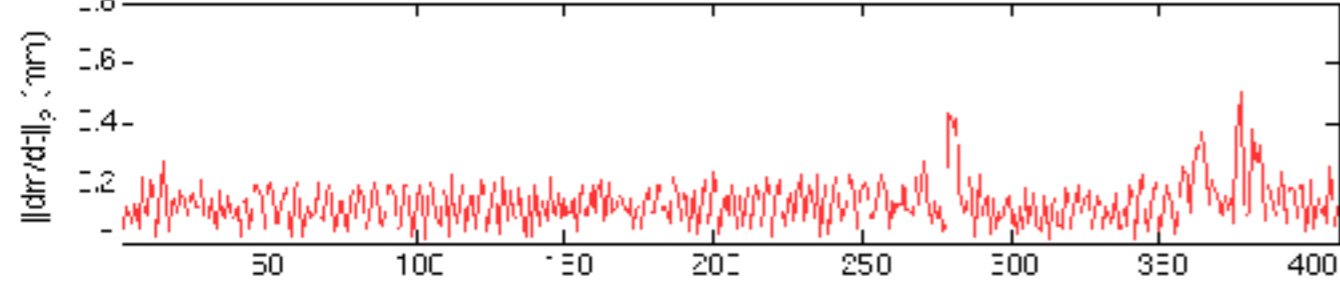
Order of eigenvalues  
239.0207@16-th PC

Order of eigenvalues  
9.4089% with 16PCs

Order of eigenvalues

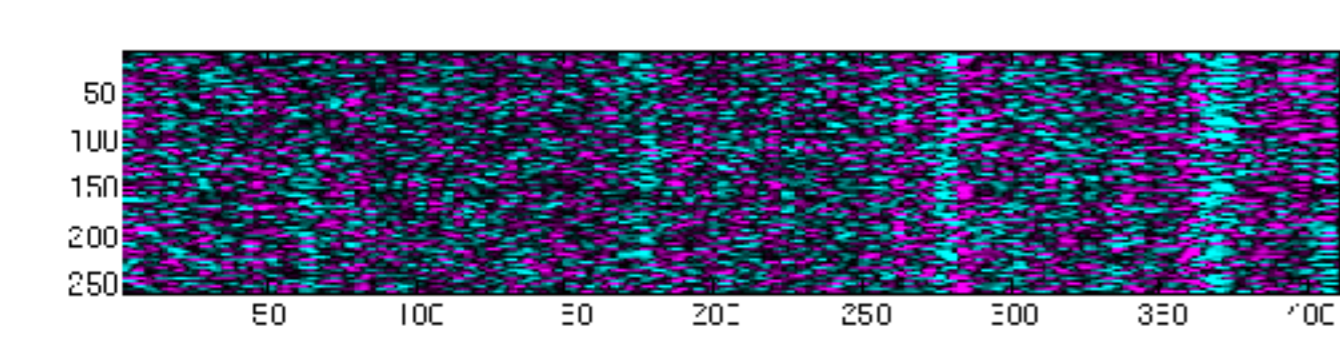


200 | square=110 nll:wmcsf0.99\_n16d | v | b | 2.00-Inf\_3.0std\_0.5mm\_b0.0 | -0.1orlg+td+rp-cc+ps+sc+cs

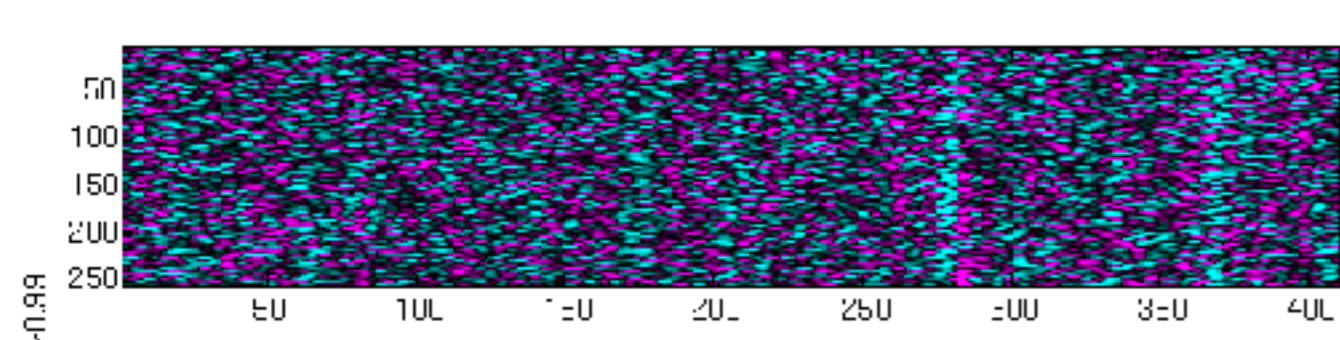


$\hat{b}_{gray}$

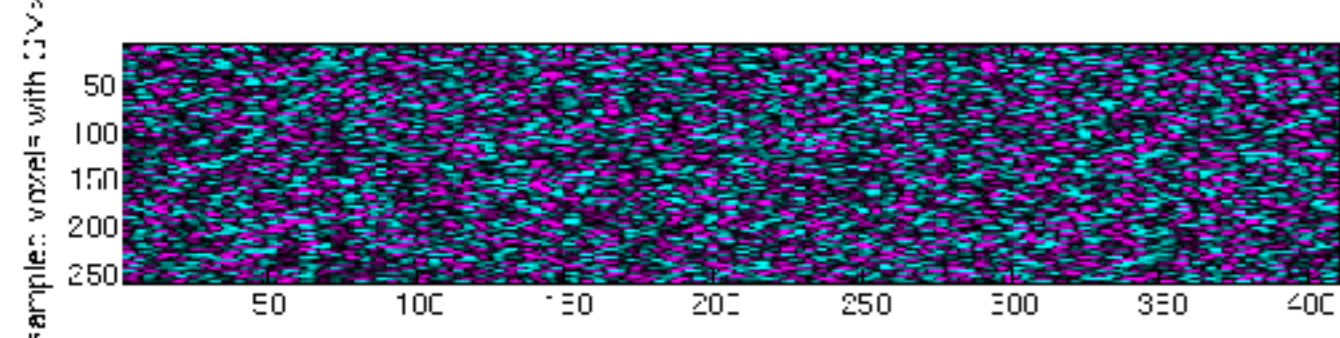
Head motion  
(frame-wise)



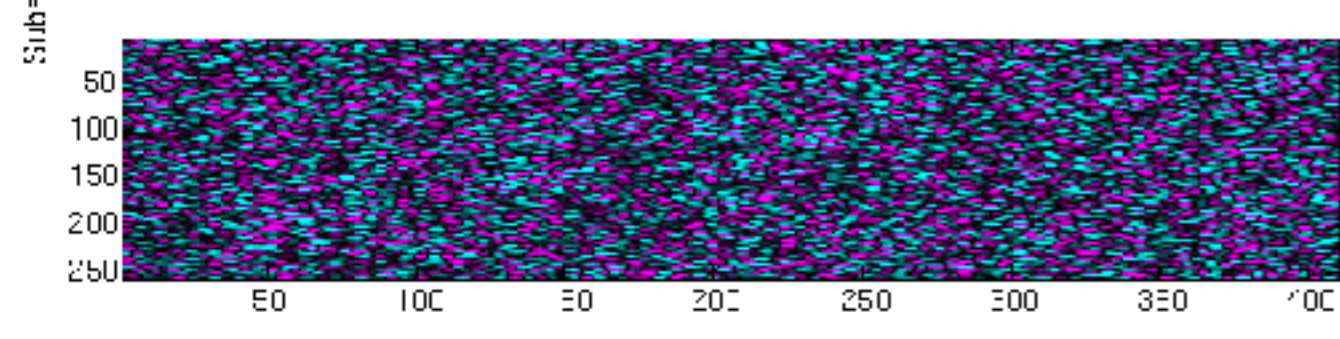
Original



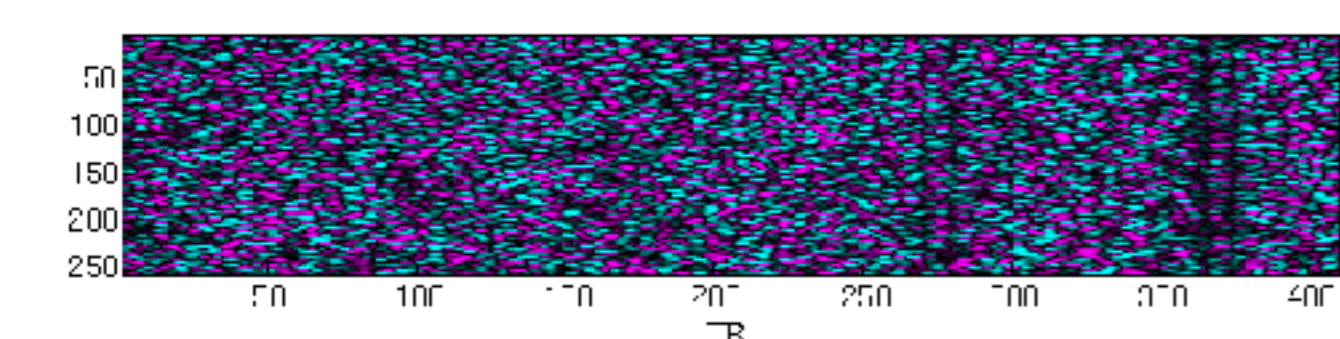
detrending (2)  
+ rigidmotion (6+1)



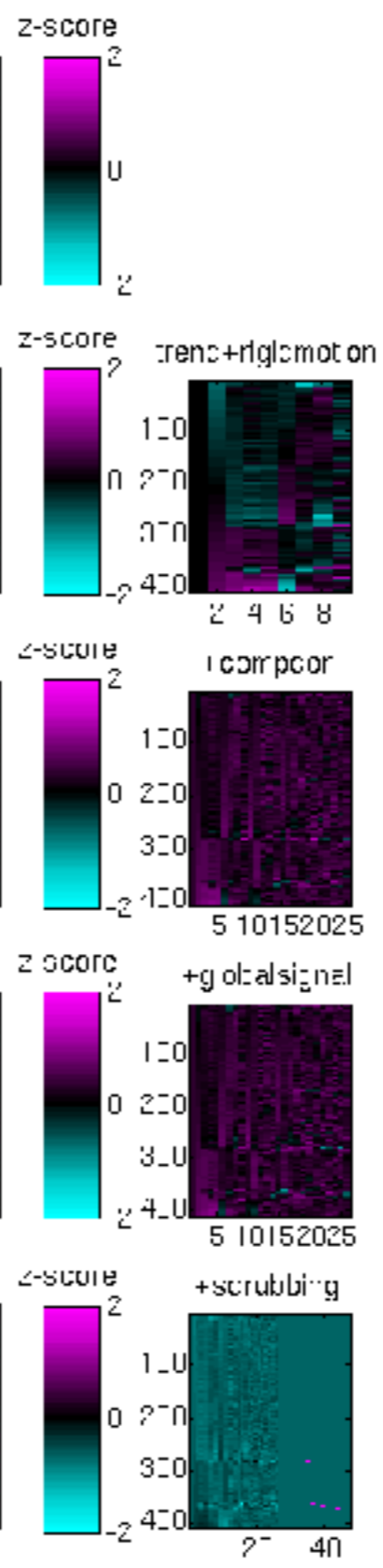
+ CompCor  
(16 PCs)



+ Global signal

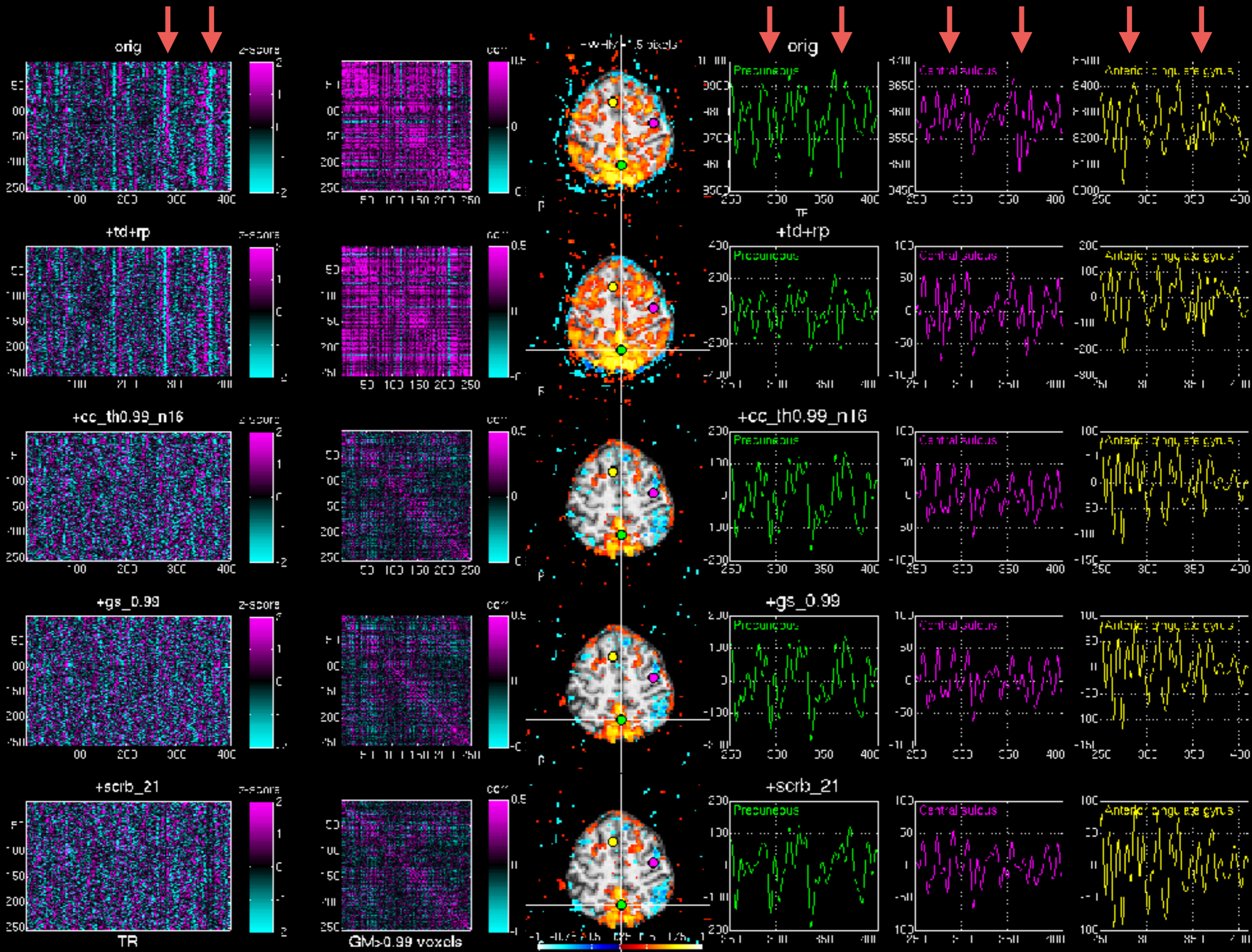


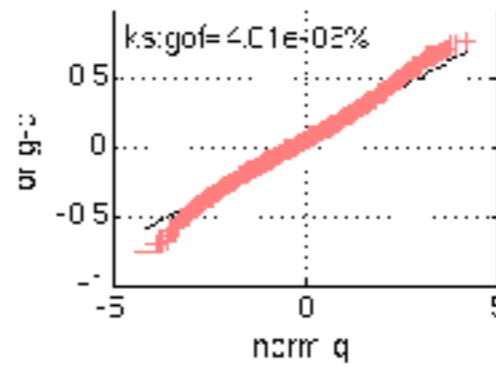
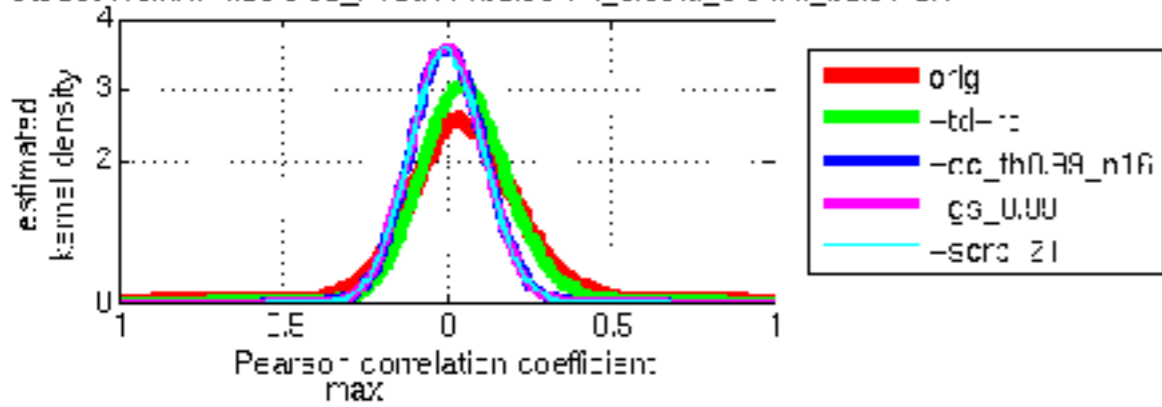
+ Scrubbing  
(21 outliers)



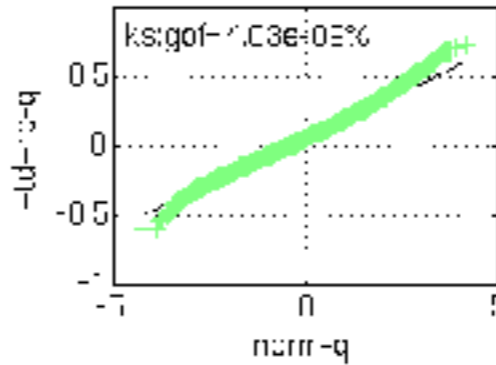
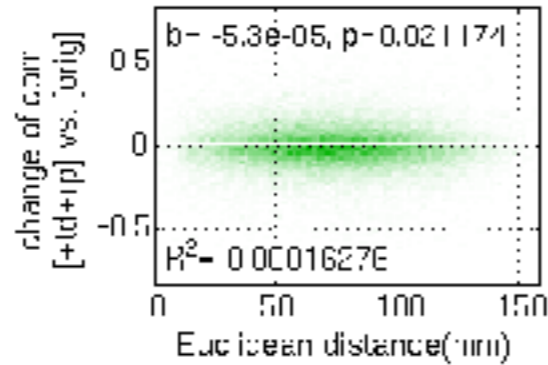
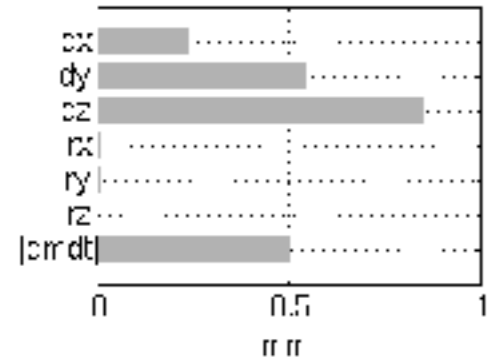


GM>0.99 voxels in the slice

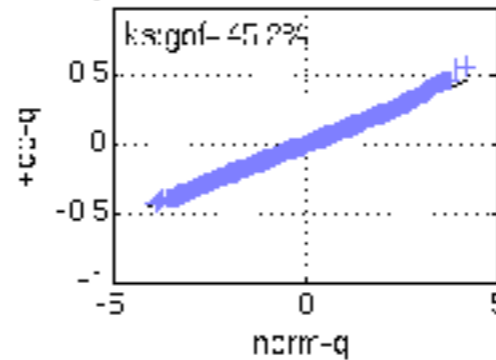
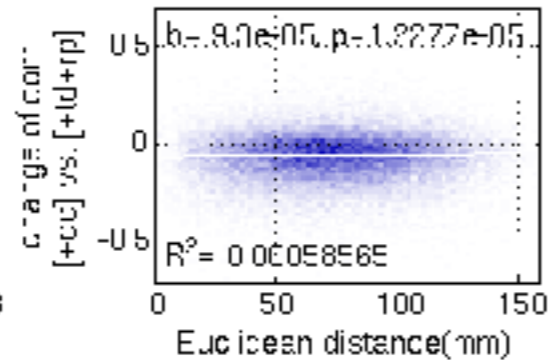
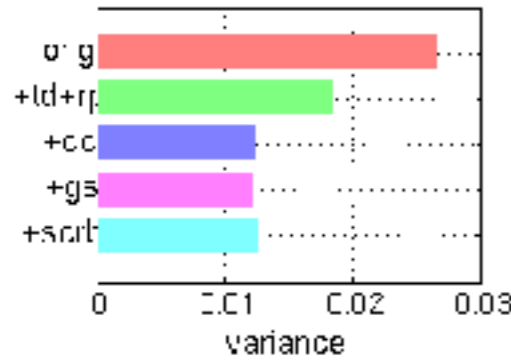




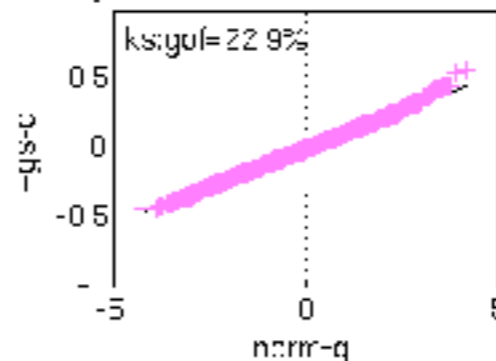
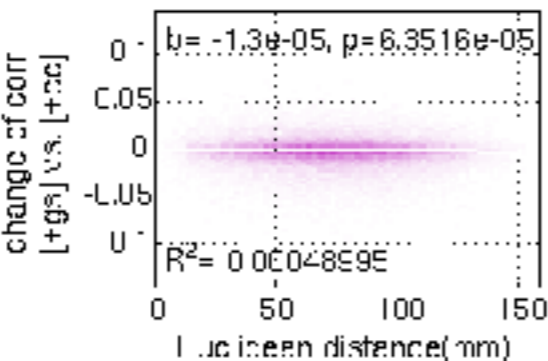
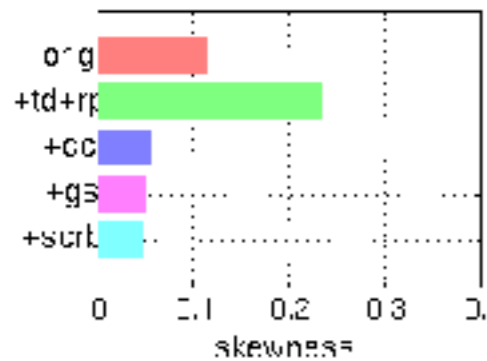
Original



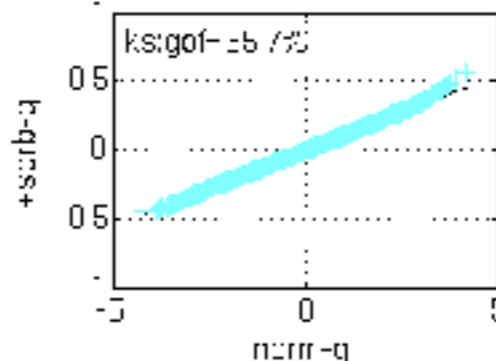
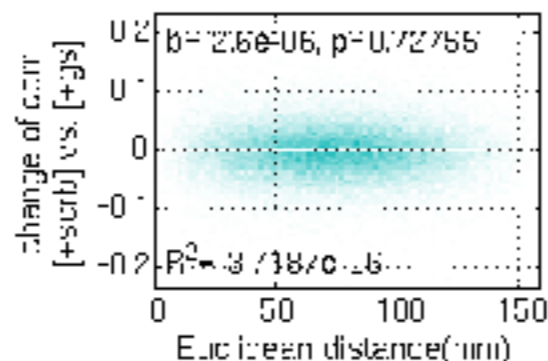
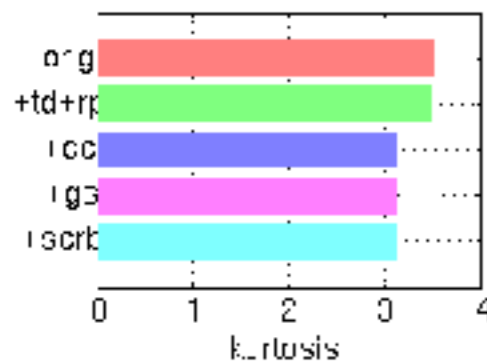
+ detrending (2)  
+ rigidmotion (6+1)



+ CompCor  
(16 PCs)



+ Global signal



+ Scrubbing  
(21 outliers)

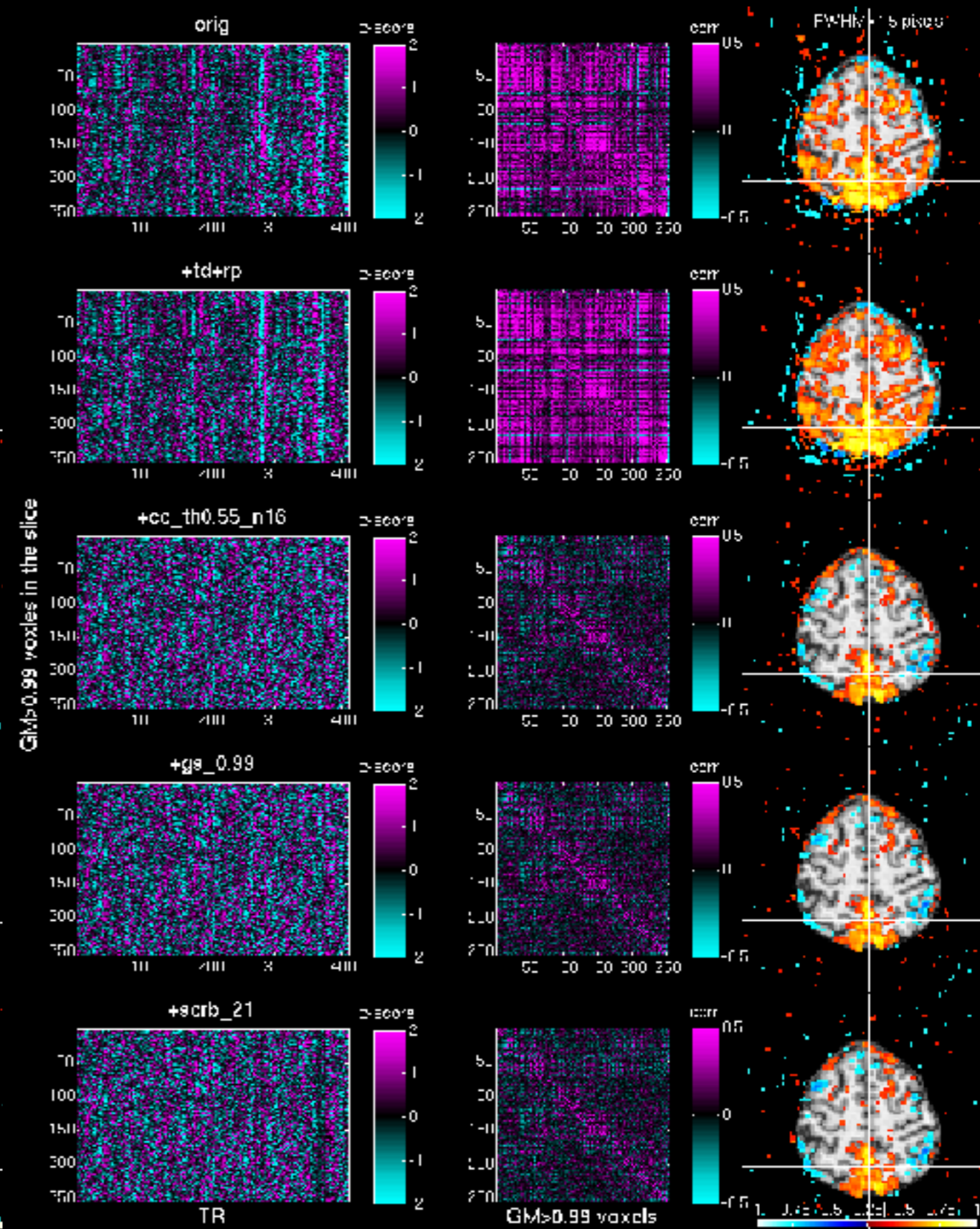
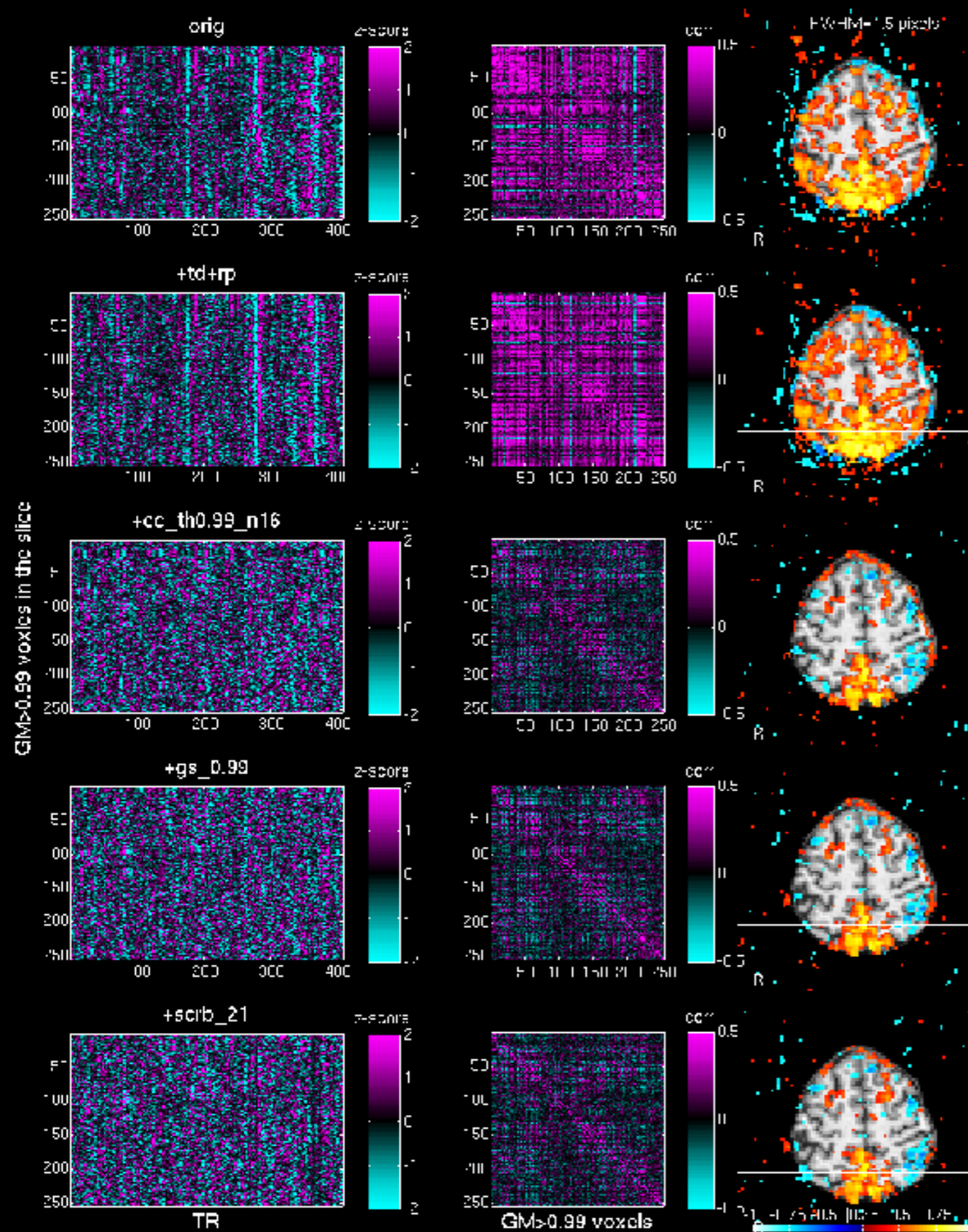
# Question #3-1

- When extracting CompCor regressors from 'WM' and 'CSF' voxels, does WM/CSF threshold matter?
- Tissue probability  $> 0.99$  vs.  $>0.55$



# WM/CSF > 0.99

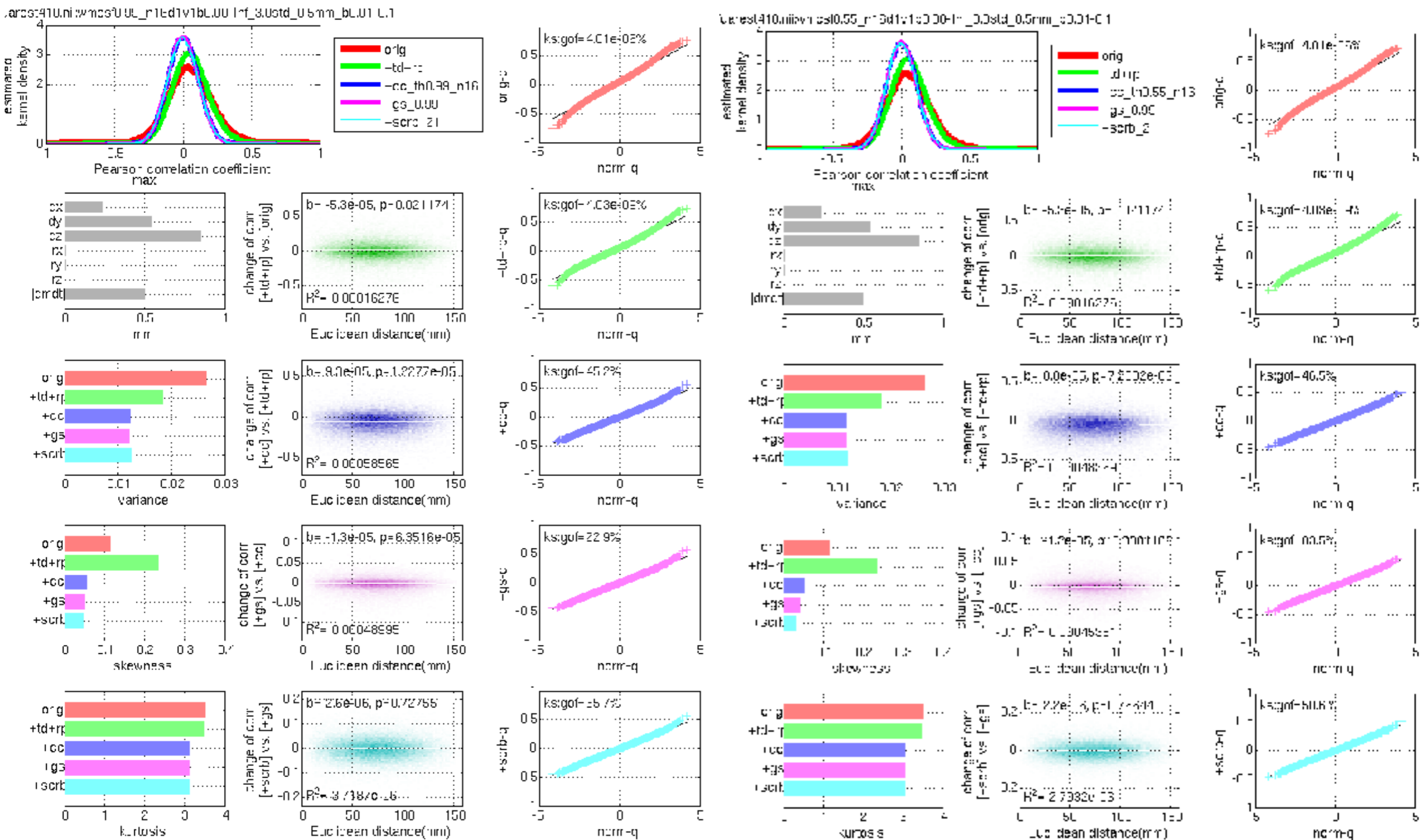
# WM/CSF > 0.55





WM/CSF > 0.99

WM/CSF > 0.55



Doesn't matter without *invasive* regressors (gs, scrubbing)

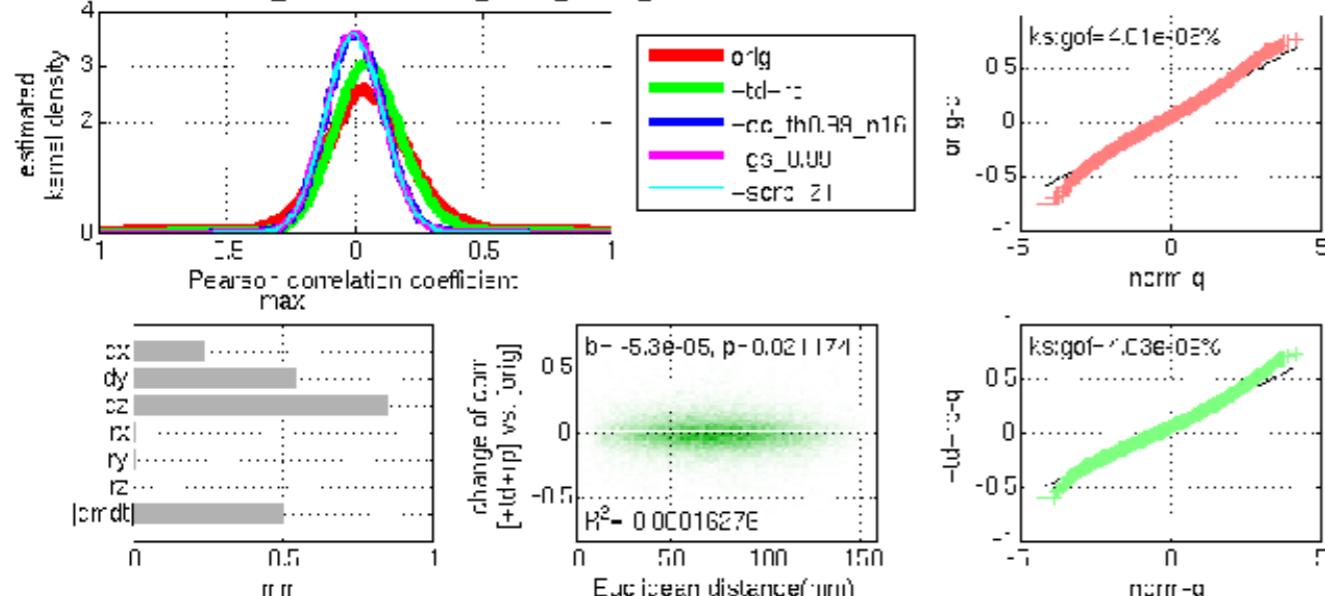
# Question #3-2

- When estimating non-neuronal 'physiological noise' from the WM/CSF voxels, averaging works as well as PCA?
- Top 16 PCs vs. mean of WM/CSF

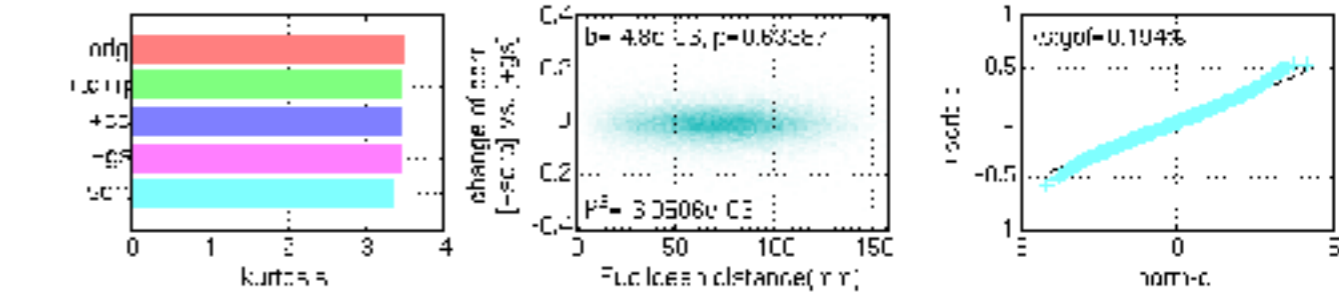
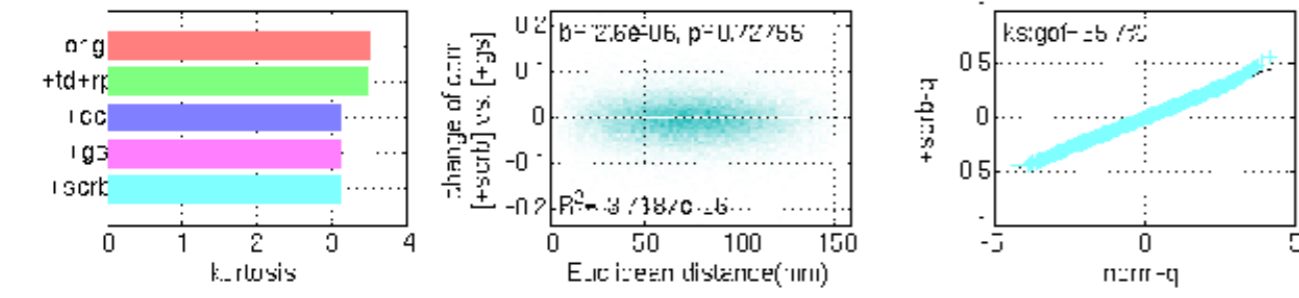
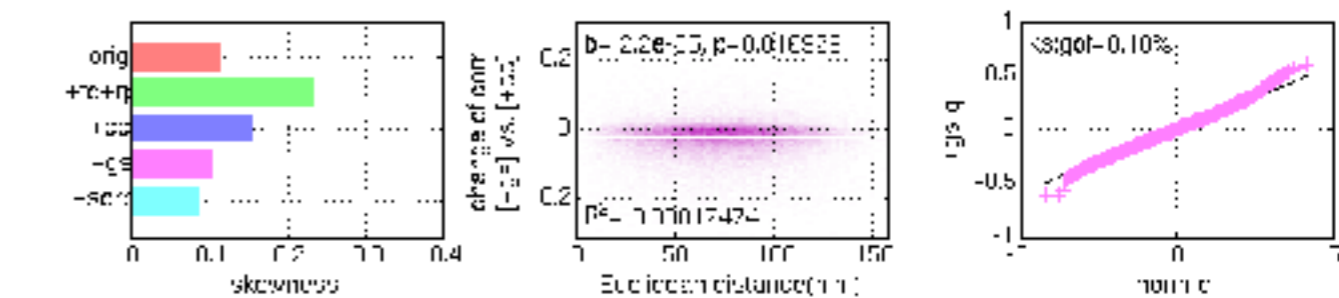
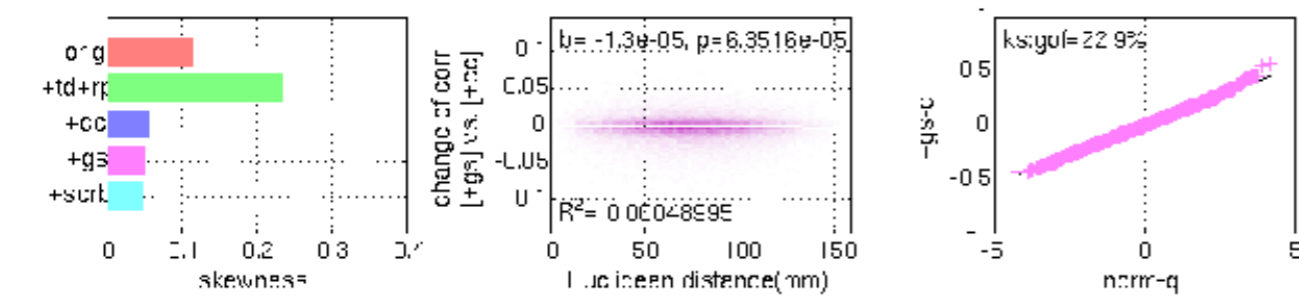
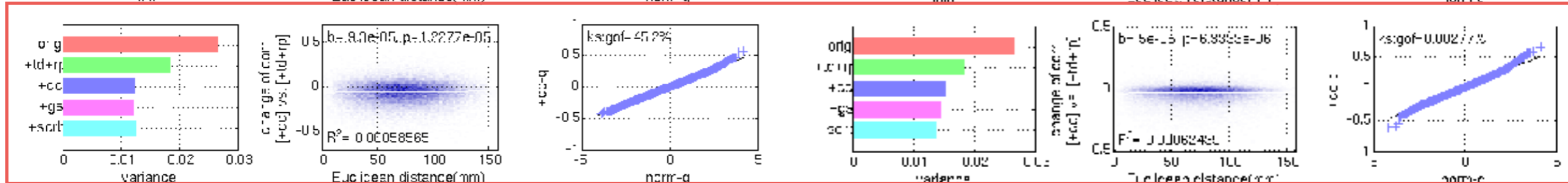
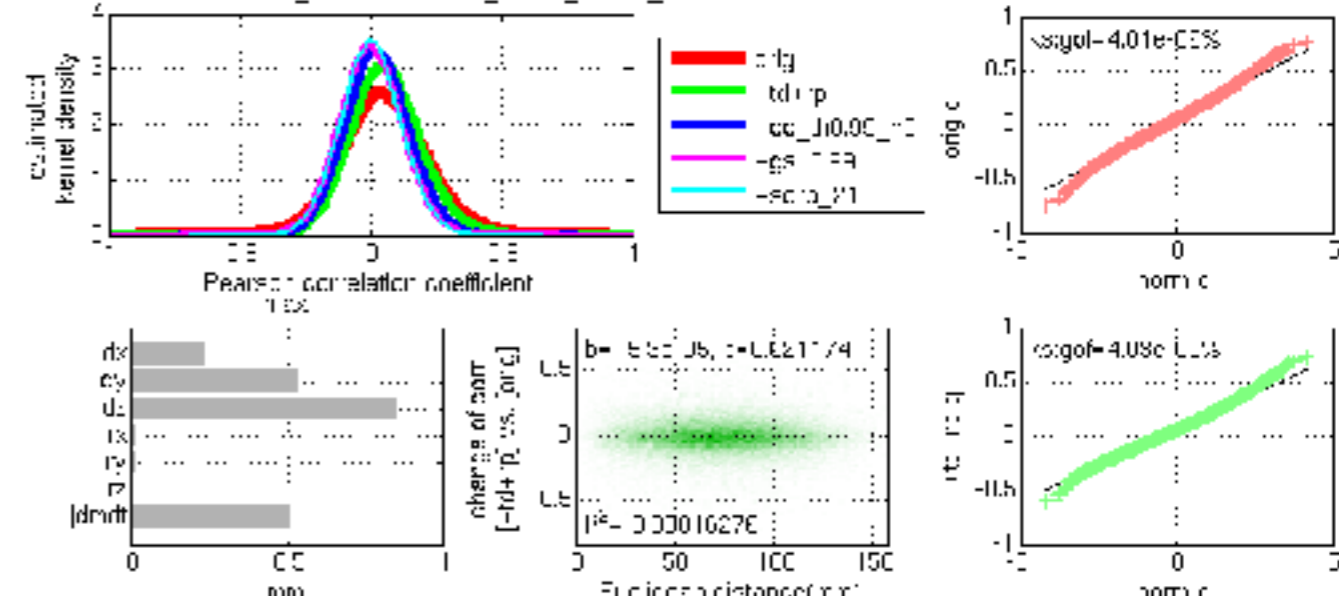
# Top 16 PCs from WM/CSF

# Mean WM/CSF

Path: /Users/ni/mcs/UUL\_r1Ed1v1BLUU/Inf\_3.Ustd\_U5mm\_bLU1/L1



Path: /Users/ni/mcs/UUL\_r1Ed1v1BLUU/Inf\_3.Ustd\_U5mm\_bLU1/L1

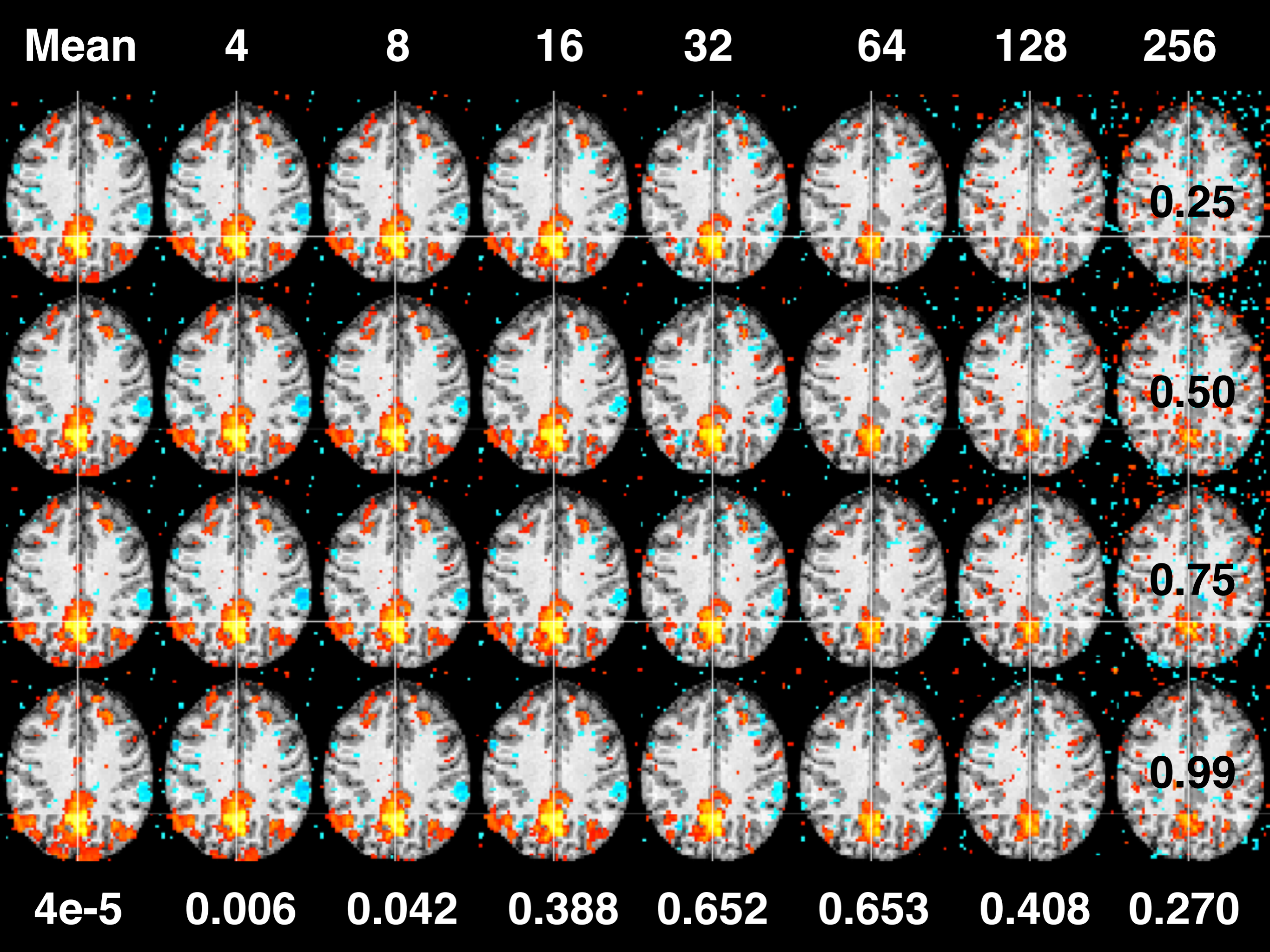


Both are better than just rigidmotion parameters

# Question #3

- So which regressors and parameters should I use?
  - WM/CSF threshold
  - # of CompCor regressors
  - Global signal? Scrubbing?
  - Different combinations of regressors?





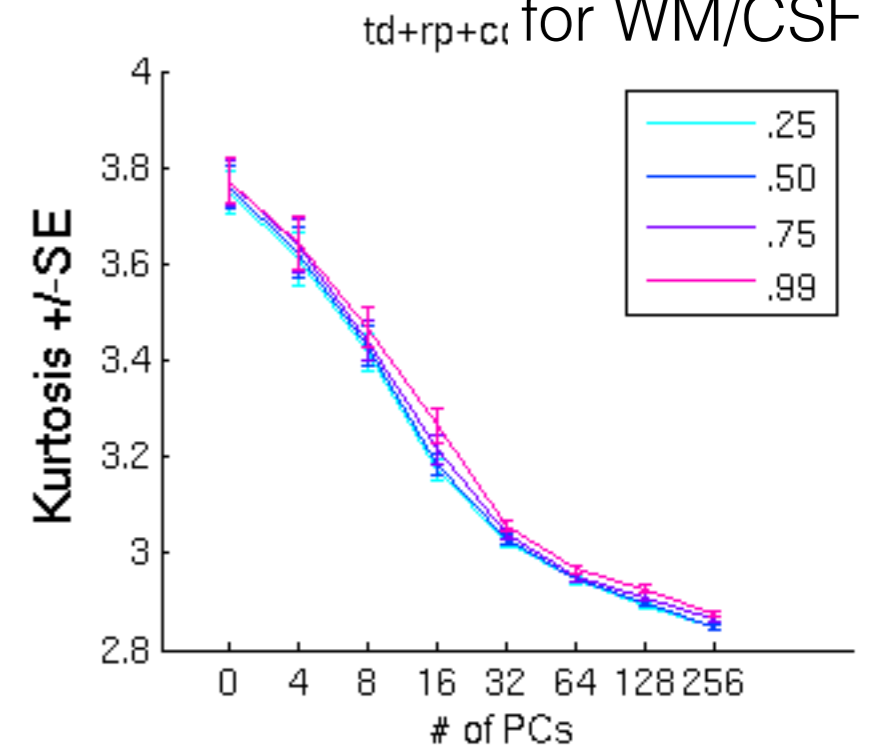
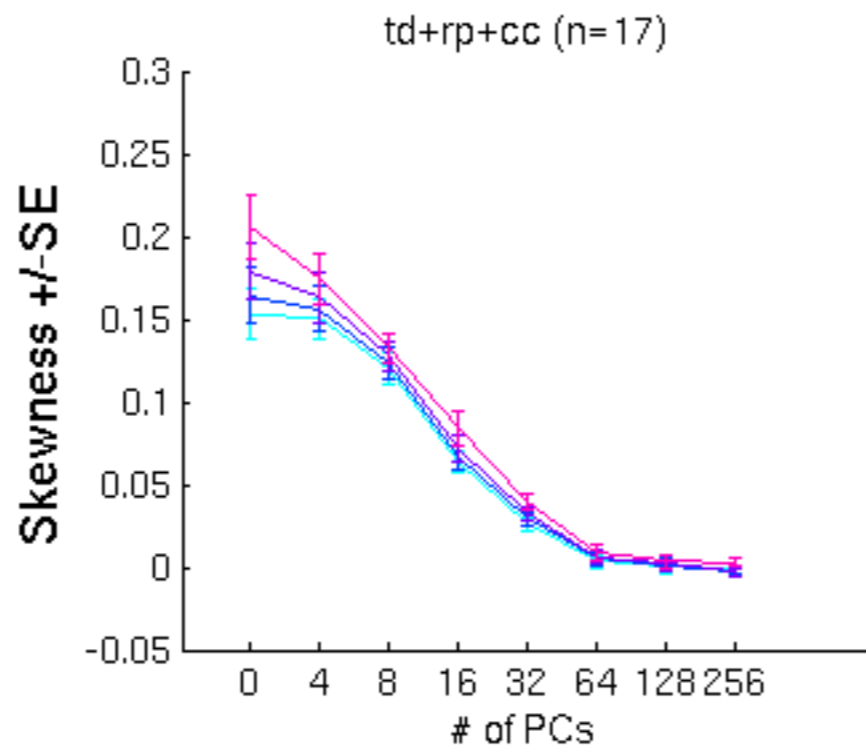
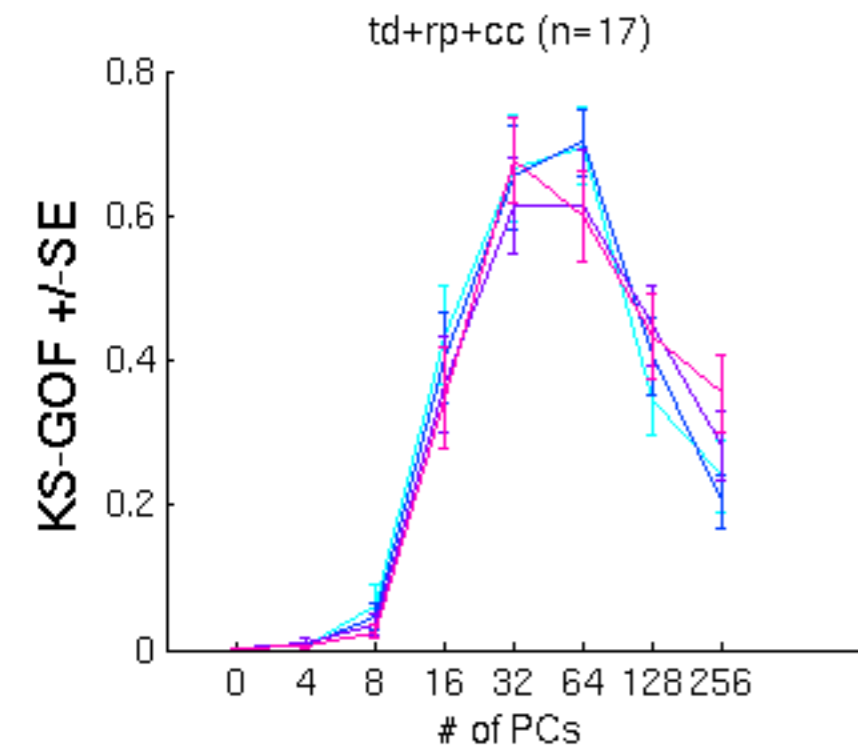




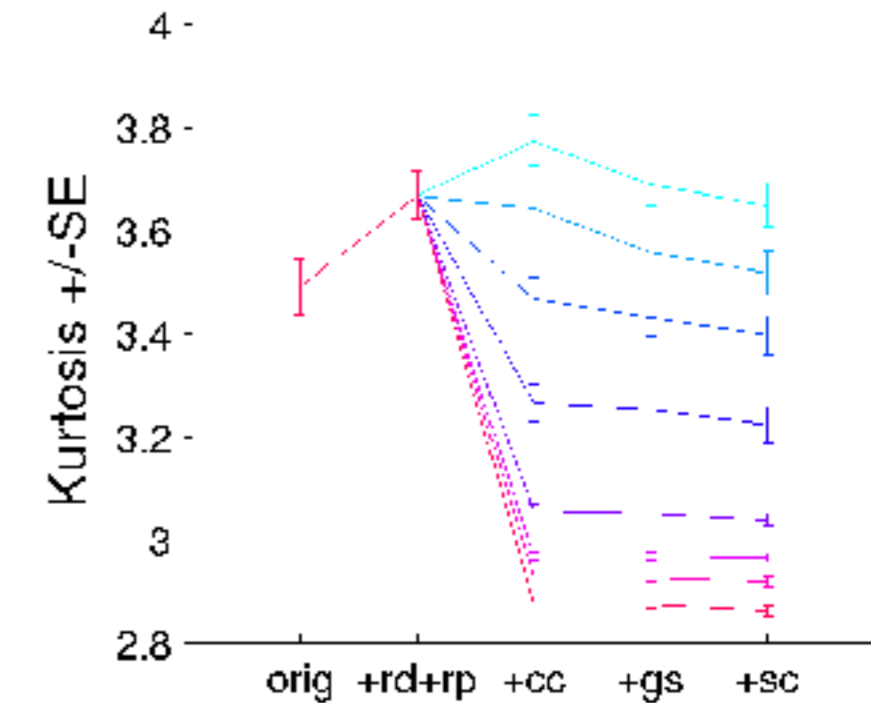
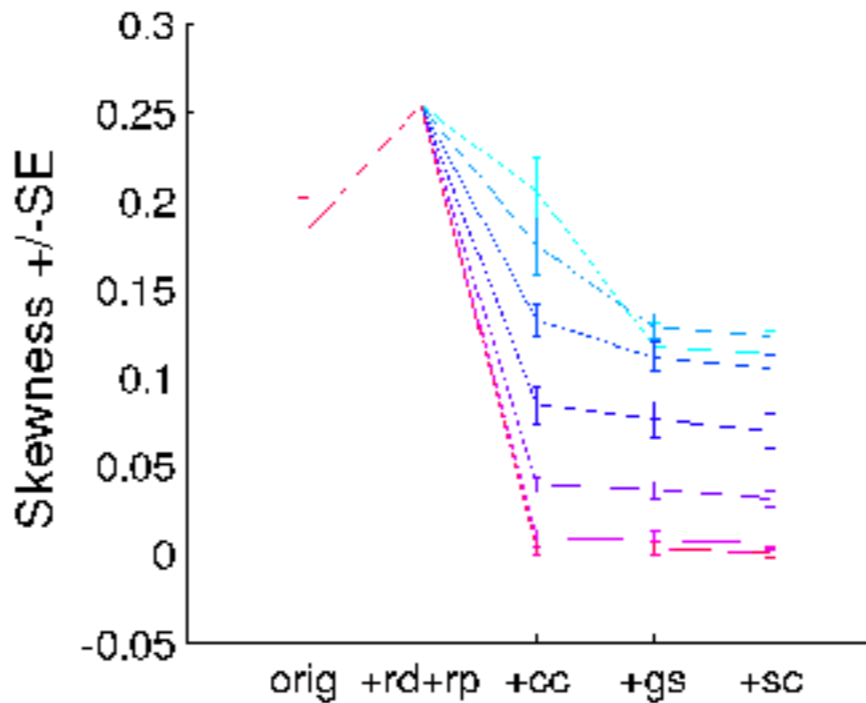
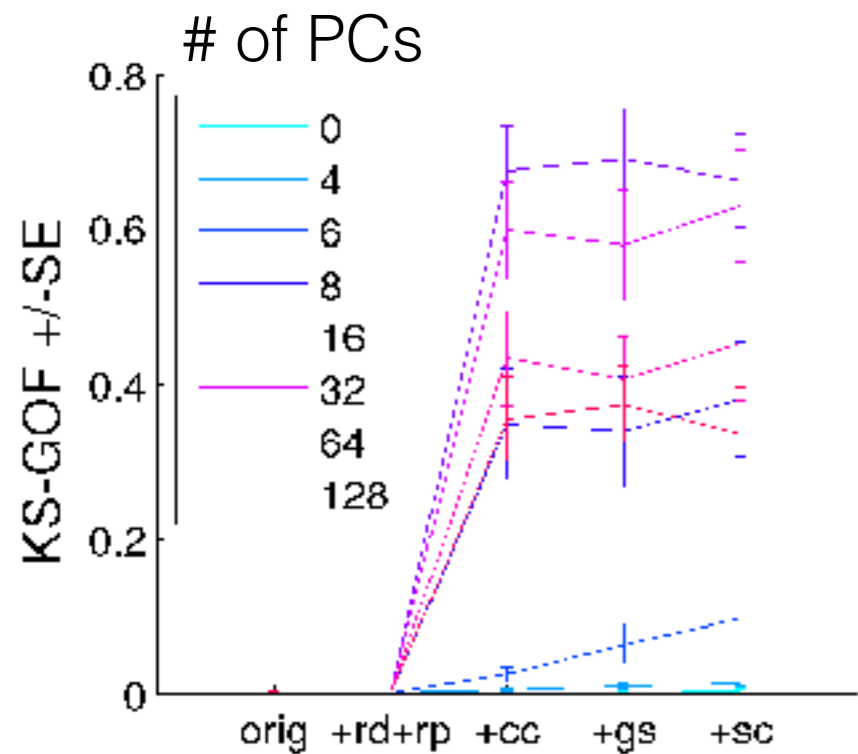


$b = 1 + \text{trend} + \text{motion} + \text{CompCor} (n=17)$

Threshold  
for WM/CSF



WM/CSF > 0.99 (n=17)



# Conclusion

- Head motion spuriously heightens correlation between BOLD timeseries (directly affects topological measures such as degree centrality).
- CompCor regressors, or at least the mean, from WM/CSF voxels normalize correlation distribution.
- Denoising regressors should be tailored to the nature of the data on hands (e.g., a children study may benefit from scrubbing).

And further  
discussion!