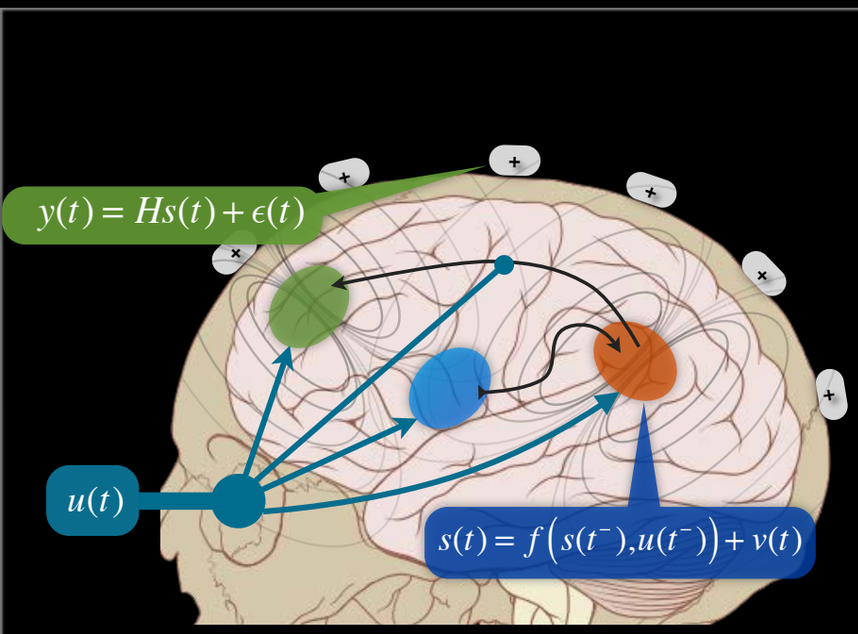
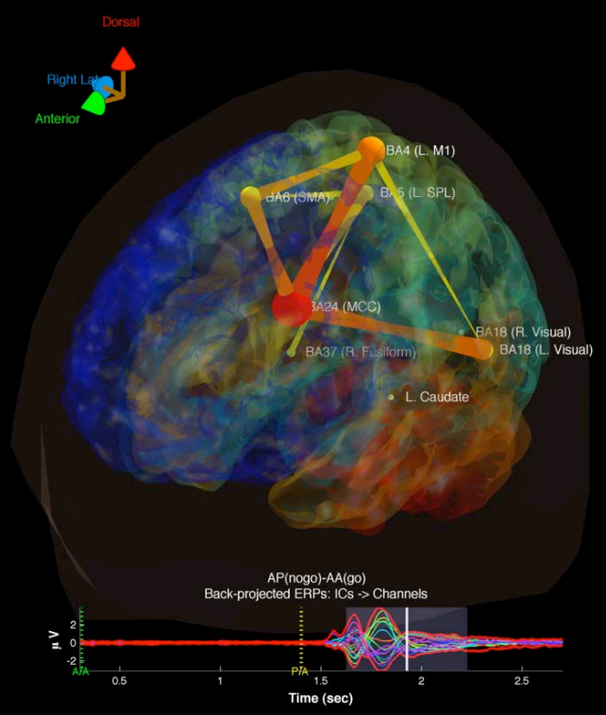
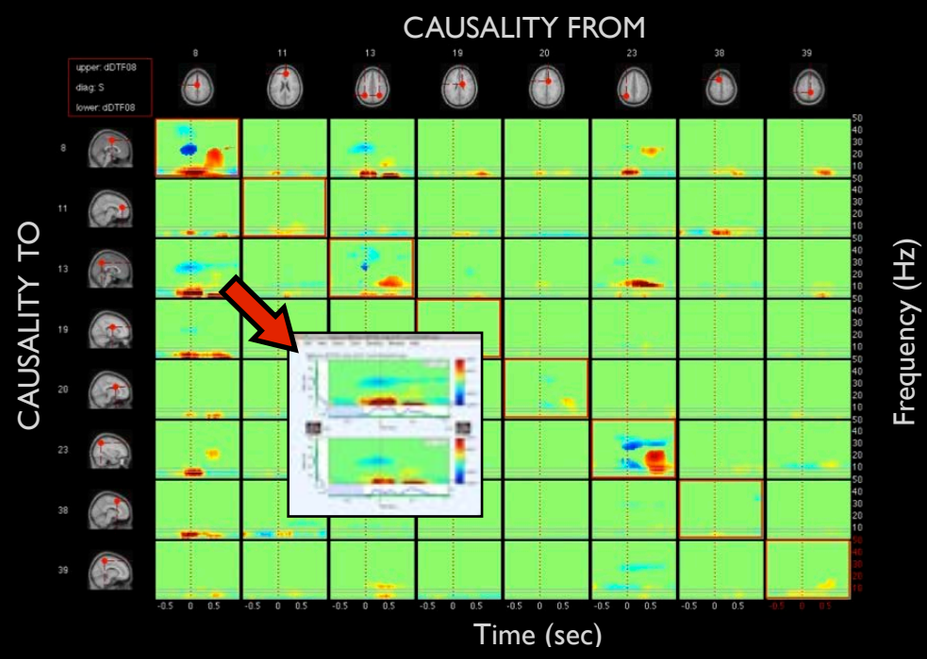
The Dynamic Brain II: Modeling Neural Dynamics and Interactions from M/EEG







Tim Mullen, PhD

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Review

Theoretical Foundations I

- Functional Connectivity Measures (PLV, PAC, Coherence)
- Effective Connectivity Measures and Granger Causality
- Scalp versus Source
- Adapting to Time-Varying Dynamics

Practicum: Hands-On Walkthrough of SIFT

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Outline

Theoretical Foundations II

- Model Validation
- Multivariate vs. Bivariate
- Imposing Constraints
- Single-trial Estimation and State-Space Models
- Statistical Testing

Practicum: Hands-On Simulation-based training

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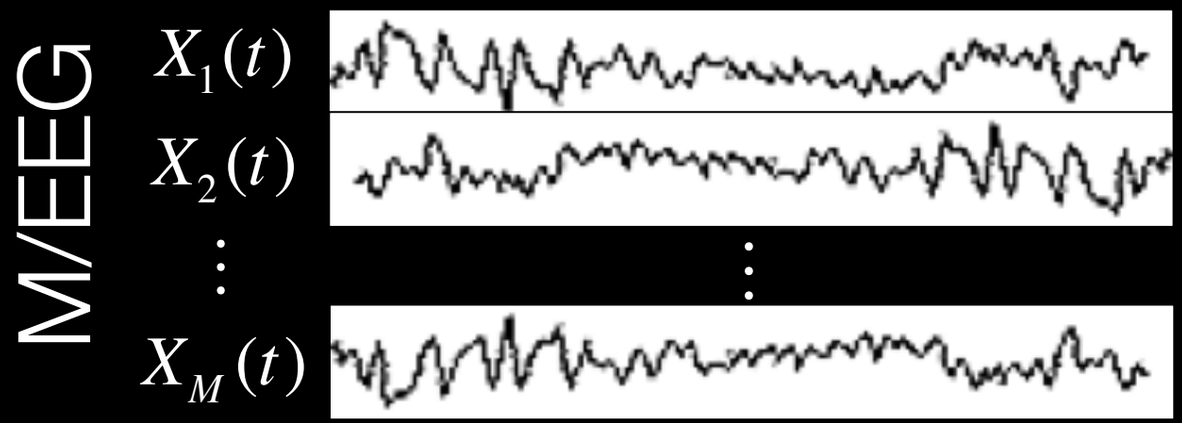
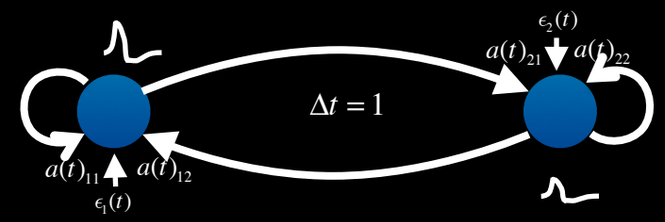
Review: The VAR model

- The Vector Autoregressive (VAR) model as a basis for dynamical estimation



Stochastic Linear Dynamical System

$$X_1(t) = a(t)_{11} X_1(t-1) + a(t)_{12} X_2(t-1) + \epsilon_1(t)$$
$$X_2(t) = a(t)_{22} X_2(t-1) + a(t)_{21} X_1(t-1) + \epsilon_2(t)$$



VAR

$$\mathbf{X}(t) = \sum_{k=1}^{p} \mathbf{A}^{(k)}(t)\mathbf{X}(t-k) + \mathbf{E}(t)$$

Granger Causality | Coherence | Spectrum | ...

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Model Validation

- If a model is poorly fit to data, then few, if any, inferences can be validly drawn from the model.
- There a number of criteria which we can use to determine whether we have appropriately fit our VAR model. Here are three commonly used categories of tests:
- **Whiteness Tests:** checking the residuals of the model for serial and cross-correlation
- **Consistency Test:** testing whether the model generates data with same correlation structure as the real data
- **Stability Test:** checking the stability/stationarity of the model.

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Whiteness Tests

- We can regard the VAR[p] model coefficients $\mathbf{A}^{(k)}$ as a filter which transforms random (white) noise $\mathbf{E}(t)$, into observed, structured data $\mathbf{X}(t)$:

$$\mathbf{X}(t) = f(L)\mathbf{E}(t)\ , \quad f(L) = \left(I - \sum_{k=1}^{p} A^{(k)} L^k\right)^{-1}$$

L is a "lag operator"

$$L^k Z(t) = Z(t-k)$$

- For coefficient estimates $\hat{\mathbf{A}}^{(k)}$, we can obtain the residuals

$$\hat{\mathbf{E}}(t) = \mathbf{X}(t) - \sum_{k=1}^{p} \hat{\mathbf{A}}^{(k)}(t)\mathbf{X}(t-k)$$

- If we have adequately modeled the data, the residuals should be indistinguishable from a white noise process. Correlation structure in the residuals means there is still correlation structure in the data that has not been explained by the model.

- Checking the whiteness of residuals typically involves testing whether the residual **auto-** and **cross-correlation** coefficients up to some desired lag *h* are sufficiently small to ensure that we cannot reject the null hypothesis of white residuals at some desired significance level.

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Whiteness Tests

$$\mathbf{E}(t) = N(\mathbf{0}, \mathbf{V})$$

$C_l = \left\langle \hat{\mathbf{E}}(t)\hat{\mathbf{E}}'(t-l) \right\rangle$ autocovariance at lag *l* ...

$R_l = D^{-1} C_l D^{-1}$ with correponding autocorrelation *R*

$D = diag\left(\sqrt{diag(C_0)}\right)$

$R_h = (R_1, \ldots, R_h)$ set of autocorrelations up to lag *h*

We want to test the null hypothesis $H_0 : \mathbf{R}_h = (R_1, \ldots, R_h) = 0$

against the alternative: $H_1 : \mathbf{R}_h \neq 0$

Two possible ways to do this:

- Autocorrelation function test
- Portmanteau tests

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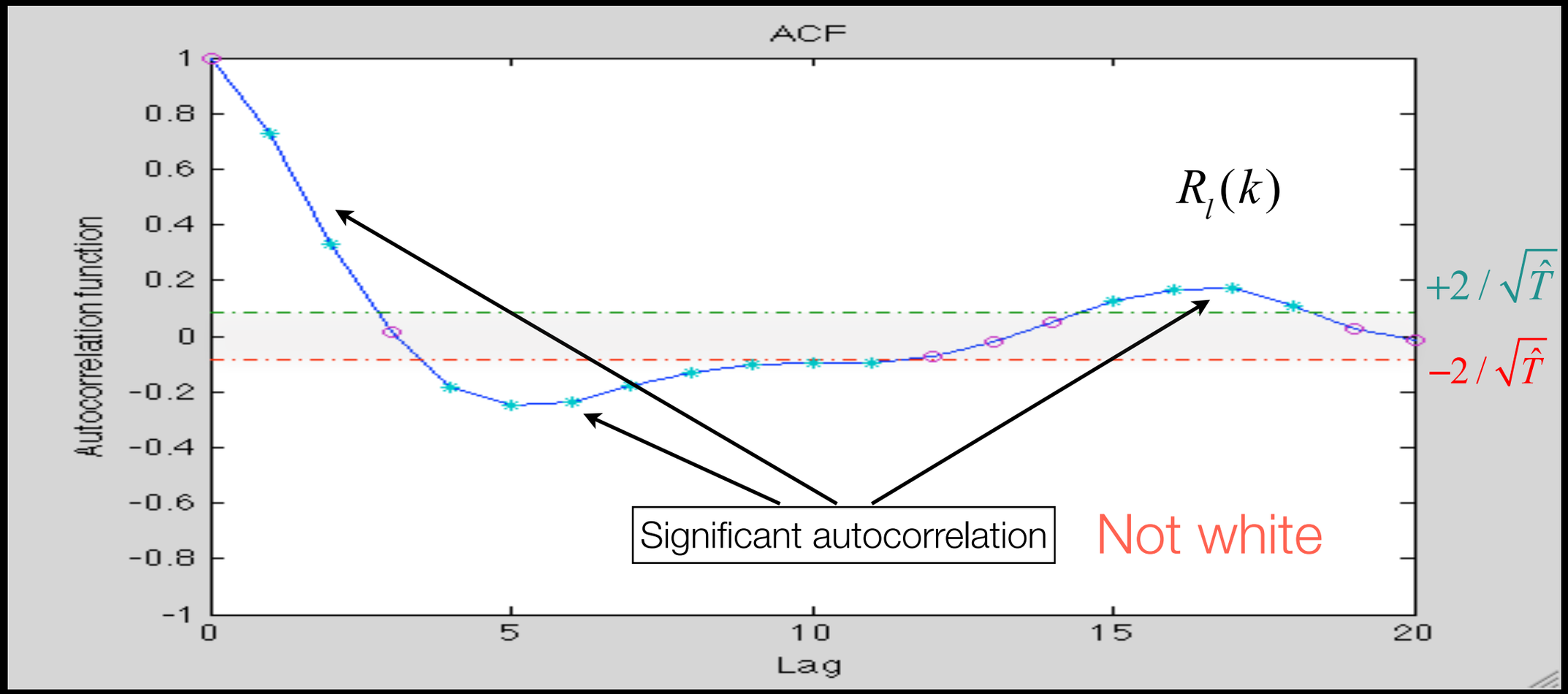
Whiteness Tests: ACF



Under the null hypothesis that $\hat{\mathbf{E}}(t)$ is Gaussian white noise, we expect approximately 1/20=5% of a.c.f. coefficients to exceed the threshold $\pm 2/\sqrt{\hat{T}}$ This gives us a p-value for rejecting H_0

$$\rho = \frac{\text{count}\left(\left|\mathbf{R}_h\right| > 2/\sqrt{\hat{T}}\right)}{\text{count}\left(\mathbf{R}_h\right)} = \frac{\text{count}\left(\left|\mathbf{R}_h\right| > 2/\sqrt{\hat{T}}\right)}{M^2(h+1) - M}$$

If $p<0.05$ ($1\text{-}p > .95$) then we cannot reject H_0 at the 5% level and we accept that residuals $\hat{\mathbf{E}}(t)$ are white

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Whiteness Tests: ACF

- Problem:
  - Confidence intervals apply to individual coefficients and assume coefficients are asymptotically independent. This may not be the case for multivariate models.
    - In small sample conditions (small *T*), this test may cause us to reject the null hypothesis (residuals indicated as non-white) **less often** than we should for the chosen significance level (Lutkepohl, 2006) -- in other words, we may have a **higher false positive rate** for accepting that the model fits the data.
- This motivates the use of alternate multivariate tests

Whiteness Tests: Portmanteau

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Table 3. Popular portmanteau tests for whiteness of residuals, implemented in SIFT. Here $\hat{T} = TN$ is the total number of samples used to estimate the covariance

Portmanteau Test	Formula (Test Statistic)	Notes
Box-Pierce (BPP)	$Q_h := \hat{T}\sum_{l=1}^{h} \mathrm{tr}\left(C_l' C_0^{-1} C_l\, C_0^{-1}\right)$	The original portmanteau test. Potentially overly-conservative. Poor small-sample properties.
Ljung-Box (LBP)	$Q_h := \hat{T}(\hat{T}+2)\sum_{l=1}^{h} (\hat{T}-l)^{-1} \mathrm{tr}\left(C_l' C_0^{-1} C_l\, C_0^{-1}\right)$	Modification of BPP to improve small-sample properties. Potentially inflates the variance of the test statistic. Slightly less conservative than LMP with slightly higher (but nearly identical) statistical power.
Li-McLeod (LMP)	$Q_h := \hat{T}\sum_{l=1}^{h} \mathrm{tr}\left(C_l' C_0^{-1} C_l\, C_0^{-1}\right) + \frac{M^2 h(h+1)}{2\hat{T}}$	Further modification of BPP to improve small-sample properties without variance inflation. Slightly more conservative than LBP. Probably the best choice in most conditions.

Mullen, 2010 (SIFT Manual)

These test statistics are asymptotically χ^2 distributed with $M^2(h-p)$ d.f.

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Consistency Tests

- A well-fit model should be able to generate data that has the same correlation structure as the original data.
- One test of this is *percent consistency* (Ding et al, 2000)
- Here we generate simulated data from our fitted model (feeding it white noise) and calculate auto- and cross-correlations up to a fixed lag for both simulated data ($\mathbf{R}_s$) and real data ($\mathbf{R}_r$).
- The percent consistency (PC) is then given by

$$\mathrm{PC} = \left(1 - \frac{\left\|\mathbf{R}_s - \mathbf{R}_r\right\|_2}{\left\|\mathbf{R}_r\right\|_2}\right) \times 100$$

- A PC value near 100% indicates that the model is able to generate data that has a nearly identical correlation structure as the original data. A PC value near 0% indicates a complete failure to model the data.

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Stability Tests

- **Stability**
  - All eigenvalues of the system matrix are ≤1
  - A stable process will not “blow up” (diverge to infinity)
  - A stable model is always a stationary model (however, the converse is not necessarily true). If a stable model adequately fits the data (white residuals), then the data is likewise stationary

Multivariate versus Bivariate

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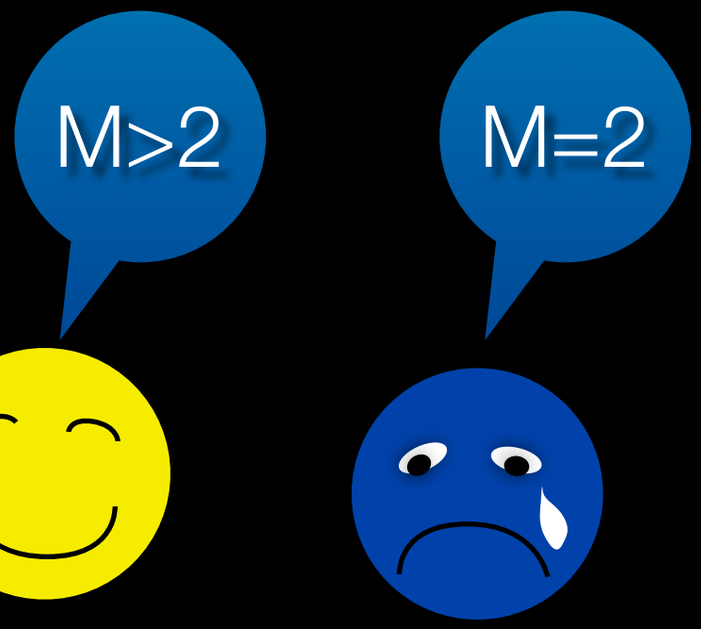
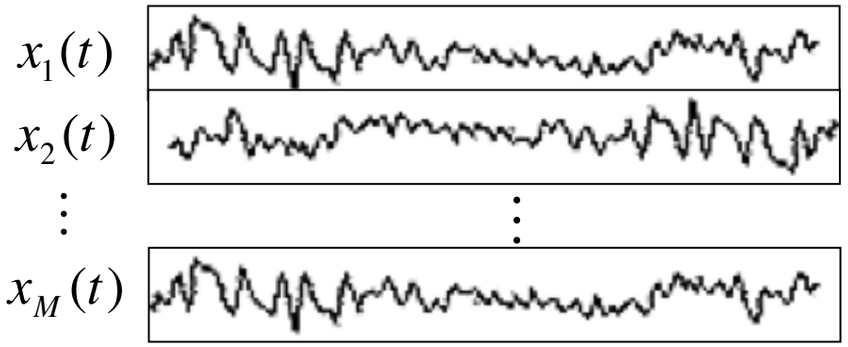
- Exclusion of processes that may exert causal influence on modeled processes increases the risk of causal mis-identification. (c.f. Pearl, *Causality: Models, Inference and Reasoning,* 2009)
  - Multivariate approaches are generally superior to bivariate approaches
    - allow detection of direct versus indirect dependence, reducing false positives
    - allow us to partially control for exogenous/unobserved causes (e.g. Guo, et al., J. Neuro. Methods, 2008)



- In the absence of *a priori* knowledge concerning causal structure, it is advisable to include as many processes as possible in a causal model *(within data/modeling limitations)*

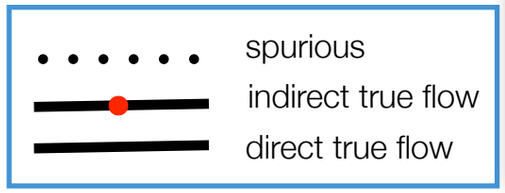


$$\mathbf{X}(t) = \sum_{k=1}^{p} \mathbf{A}^{(k)}(t)\mathbf{X}(t-k) + \mathbf{E}(t)$$

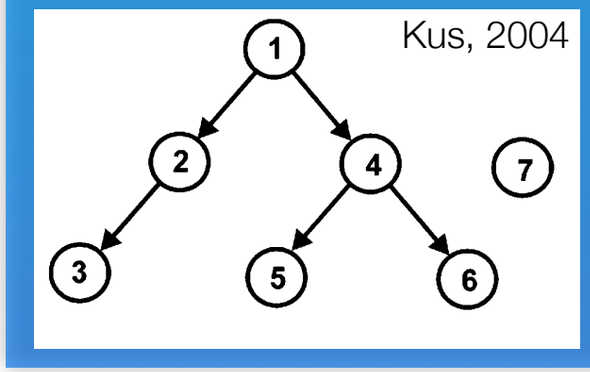
$$\mathbf{A}(f,t) = -\sum_{k=0}^{p} \mathbf{A}^{(k)}(t)e^{-i2\pi fk}; \quad \mathbf{A}^{(0)} = I$$

$$\mathbf{X}(f,t) = \mathbf{A}(f,t)^{-1}\mathbf{E}(f,t) = \mathbf{H}(f,t)\mathbf{E}(f,t)$$

NOTE: time index (t) dropped for convenience



Ground Truth

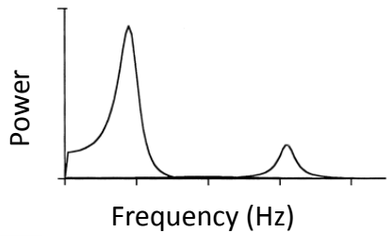


Functional

Spectrum

$$S(f) = \mathbf{X}(f)\mathbf{X}(f)^* = \mathbf{H}(f)\Sigma\mathbf{H}(f)^*$$

(Brillinger, 2001)

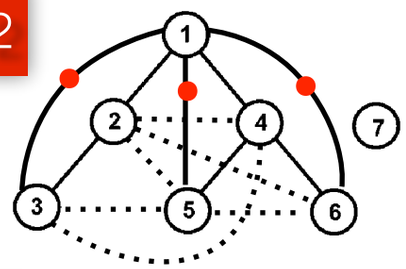


Coherency

M=2

$$C_{ij}(f) = \frac{S_{ij}(f)}{\sqrt{S_{ii}(f)S_{jj}(f)}}$$

(Brillinger, 2001)

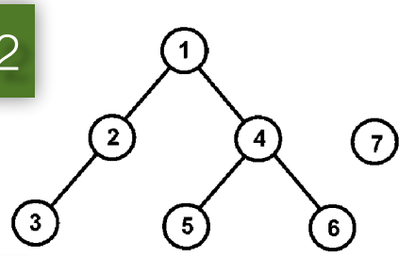


Partial Coherence

M>2

$$P_{ij}(f) = \frac{S^{-1}{}_{ij}(f)}{\sqrt{S^{-1}{}_{ii}(f)S^{-1}{}_{jj}(f}}$$

(Brillinger, 2001)



Multiple Coherence

M=1

$$G_i(f) = \sqrt{1 - \frac{\det(S(f))}{S_{ii}(f)\mathbf{M}_{ii}(f)}}$$

(Brillinger, 2001)

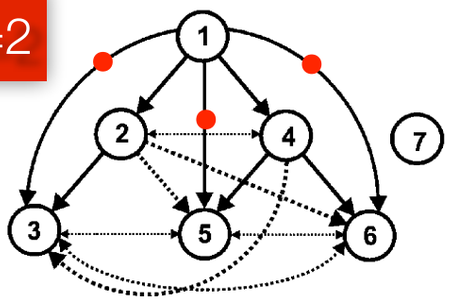
Effective

Granger-Geweke Causality

M=2

$$F_{ij}(f) = \frac{\Sigma_{jj} - (\Sigma^2{}_{ij} / \Sigma_{ii}))\,|\,H_{ij}(f)\,|^2}{S_{ii}(f)}$$

(Geweke, 1982; Bressler et al., 2007)

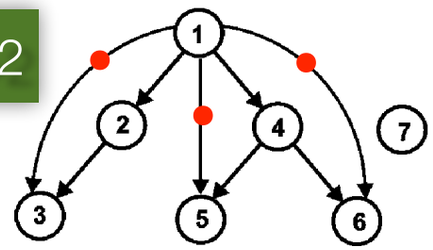


Directed Transfer Function

M>2

$$\eta^2_{ij}(f) = \frac{\left|H_{ij}(f)\right|^2}{\sum_f \sum_{k=1}^{M} \left|H_{ik}(f)\right|^2}$$

(Kaminski and Blinowska, 1991)

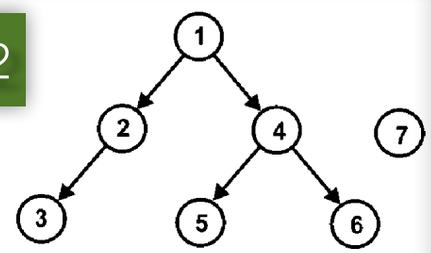


Partial Directed Coherence

M>2

$$\pi^2_{ij}(f) = \frac{|\,A_{ij}(f)\,|^2}{\sum_{k=1}^{M} \left|A_{kj}(f)\right|^2}$$

(Baccalá and Sameshima, 2001)

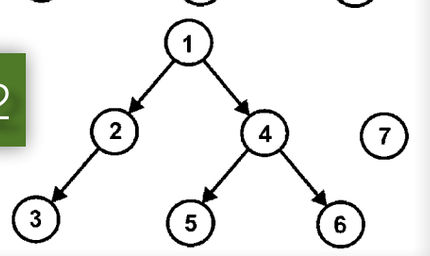


Direct DTF

M>2

$$\delta^2_{ij}(f) = \eta^2_{ij}(f)P^2_{ij}(f)$$

(Korzeniewska, 2003)



	Estimator	Formula
Spectral M.	Spectral Density Matrix	$S(f) = X(f)X(f)^* = H(f)\Sigma H(f)^*$
Coherence Measures	Coherency	$C_{ij}(f) = \frac{S_{ij}(f)}{\sqrt{S_{ii}(f)S_{jj}(f)}}$ $0 \le \lvert C_{ij}(f)\rvert^2 \le 1$
	Imaginary Coherence (iCoh)	$iCoh_{ij}(f) = \mathrm{Im}(C_{ij}(f))$
	Partial Coherence (pCoh)	$P_{ij}(f) = \frac{\hat{S}_{ij}(f)}{\sqrt{\hat{S}_{ii}(f)\hat{S}_{jj}(f)}}$ $\hat{S}(f) = S(f)^{-1}$ $0 \le \lvert P_{ij}(f)\rvert^2 \le 1$
	Multiple Coherence (mCoh)	$G_i(f) = \sqrt{1 - \frac{\det(S(f))}{S_{ii}(f)\mathbf{M}_{ii}(f)}}$ $\mathbf{M}_{ii}(f)$ is the **minor** of $S(f)$ obtained by removing the i[th] row and column of $S(f)$ and returning the determinant.

	Estimator	Formula
Partial Directed Coherence Measures	Normalized Partial Directed Coherence (PDC)	$\pi_{ij}(f) = \frac{A_{ij}(f)}{\sqrt{\sum_{k=1}^{M}\lvert A_{kj}(f)\rvert^2}}$ $0 \le \lvert \pi_{ij}(f)\rvert^2 \le 1$ $\sum_{j=1}^{M}\lvert \pi_{ij}(f)\rvert^2 = 1$
	Generalized PDC (GPDC)	$\bar{\pi}_{ij}(f) = \frac{\frac{1}{\Sigma_{ii}}A_{ij}(f)}{\sqrt{\sum_{k=1}^{M}\frac{1}{\Sigma_{ii}^2}\lvert A_{kj}(f)\rvert^2}}$ $0 \le \lvert \bar{\pi}_{ij}(f)\rvert^2 \le 1$ $\sum_{j=1}^{M}\lvert \bar{\pi}_{ij}(f)\rvert^2 = 1$
	Renormalized PDC (rPDC)	$\lambda_{ij}(f) = Q_{ij}(f) * V_{ij}(f)^{-1} Q_{ij}(f)$ where $Q_{ij}(f) = \begin{pmatrix} \mathrm{Re}[A_{ij}(f)] \\ \mathrm{Im}[A_{ij}(f)] \end{pmatrix}$ and $V_{ij}(f) = \sum_{k,l=1}^{p} R_{jj}^{-1}(k,l)\Sigma_{ii} Z(2\pi f,k,l)$ $Z(\omega,k,l)$ $= \begin{pmatrix} \cos(\omega k)\cos(\omega l) & \cos(\omega k)\sin(\omega l) \\ \sin(\omega k)\cos(\omega l) & \sin(\omega k)\sin(\omega l) \end{pmatrix}$ ***R*** is the [$(Mp)^2$ x $(Mp)^2$] covariance matrix of the VAR[p] process (Lütkepohl, 2006)
Granger-Geweke	Granger-Geweke Causality (GGC)	$F_{ij}(f) = \frac{\left(\Sigma_{jj} - (\Sigma_{ij}^2 / \Sigma_{ii})\right)\lvert H_{ij}(f)\rvert^2}{S_{ii}(f)}$

	Estimator	Formula
Directed Transfer Function Measures	Normalized Directed Transfer Function (DTF)	$\gamma_{ij}(f) = \frac{H_{ij}(f)}{\sqrt{\sum_{k=1}^{M}\lvert H_{ik}(f)\rvert^2}}$ $0 \le \lvert \gamma_{ij}(f)\rvert^2 \le 1$ $\sum_{j=1}^{M}\lvert \gamma_{ij}(f)\rvert^2 = 1$
	Full-Frequency DTF (ffDTF)	$\eta_{ij}^2(f) = \frac{\lvert H_{ij}(f)\rvert^2}{\sum_f \sum_{k=1}^{M}\lvert H_{ik}(f)\rvert^2}$
	Direct DTF (dDTF)	$\delta_{ij}^2(f) = \eta_{ij}^2(f) P_{ij}^2(f)$

$$\mathbf{X}(t) = \sum_{k=1}^{p}\mathbf{A}^{(k)}(t)\mathbf{X}(t-k) + \mathbf{E}(t)$$
$$\mathbf{A}(f,t) = -\sum_{k=0}^{p}\mathbf{A}^{(k)}(t)e^{-i2\pi fk};\ \ \mathbf{A}^{(0)} = I$$
$$\mathbf{X}(f,t) = \mathbf{A}(f,t)^{-1}\mathbf{E}(f,t) = \mathbf{H}(f,t)\mathbf{E}(f,t)$$

$H(f)$ Transfer Function

$A(f)$ System Matrix

Σ Noise Covariance Matrix

Variance Stabilization

For additional details, see SIFT Handbook (sccn.ucsd.edu/wiki/SIFT)

Multivariate Models: Limitations

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- However, multivariate methods come with a cost:
  - More parameters + limited data = higher risk of over-fitting or worse yet....
  - ...the problem becomes *ill-posed* or *under-determined*. There are insufficient observations to uniquely determine a solution to the system of equations defining our model.

Multivariate Models: Limitations

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How many samples do we need?

- N = number of samples required
- M = number of variables/sources
- T = number of trials/realizations
- p = model order
- We have M^2p model coefficients to estimate. So our ordinary least-squares solution requires a *minimum* of M^2p samples.

$$N \geq M^2 p$$

- Back-of-envelope: M=20, p=10, T=1 We need 20^2x10 = 4000 samples -- 20 second epoch at sampling rate of 200Hz!

Ensemble aggregation (T > 1)?

- M=20, p=10, T=50: 4000/50 samples/trial → 20/50 = 0.4 sec epoch

Multivariate Models: Constraints

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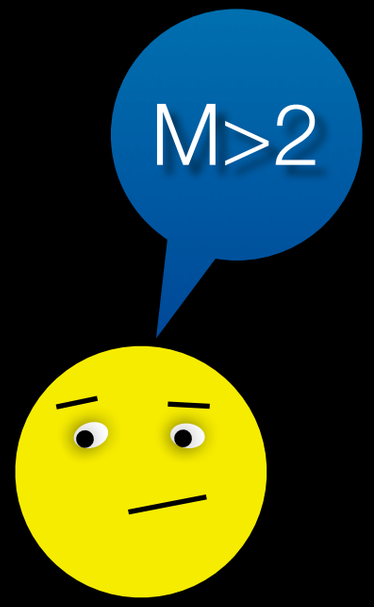

Solutions?

Make assumptions (impose constraints)

We want to *a priori* restrict the range of allowable values for our parameters -- transforming the problem from one with infinite number of solutions in the original parameter space to one with a unique (“best”) solution in the new parameter space

In a Bayesian context, this corresponds to making assumptions about the *prior distribution* of the parameters (Gaussian, Laplacian, ...)

Multivariate Models: Constraints

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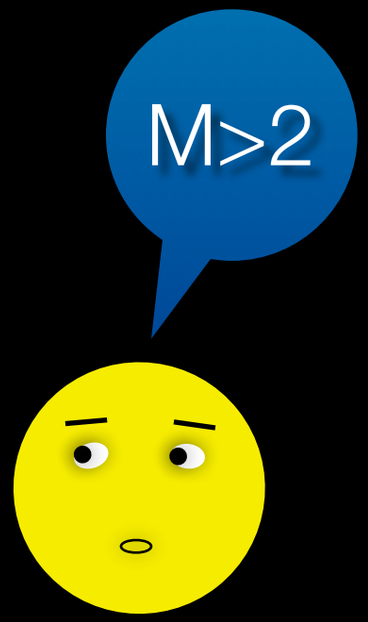
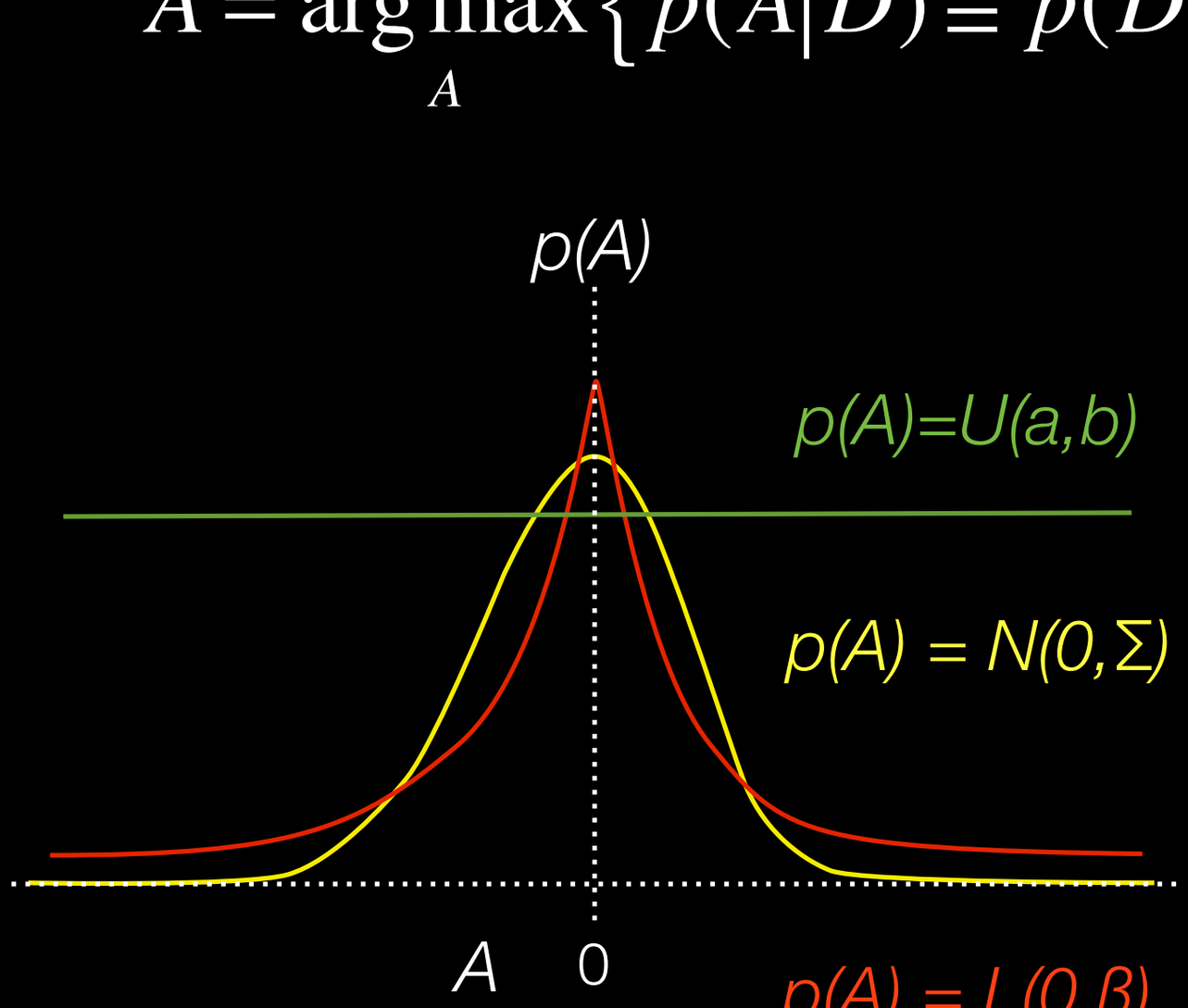
posterior | likelihood | prior

$$\hat{A} = \arg\max_{A} \left\{ p(A|D) \cong p(D|A)\,p(A) \right\}$$





Unconstrained (all values equally probable). e.g. Uniform distribution

Smoothness constraints
- large differences in values unlikely
- small (non-zero) values most probable. e.g. Normal (gaussian) prior.

Sparsity constraint
- many values small or exactly zero with occasional large values e.g. Laplacian prior

Smoothness Constraints

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- Standard least-squares solution

prediction error

$$A(t) = \arg\min_{\hat{A}} \left(\left\| Y - Z\tilde{A} \right\|_2^2 \right)$$

$$\mathbf{X}(t) = \sum_{k=1}^{p} \mathbf{A}^{(k)}(t)\mathbf{X}(t-k) + \mathbf{E}(t)$$

$$\tilde{A} = [A^{(1)}(t), \ldots, A^{(p)}(t)]^T$$

$$X_k = [X(p+1-k), \ldots, X(N-k)]^T$$

$$Z = [X_1, \ldots, X_p]$$

$$Y = X_0$$

Rewrite VAR[p] as VAR[1]

Smoothness Constraints

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- Ridge Regression
  (Tikhonov Regularization, Minimum-(L_2)-Norm

prediction error

penalty term, enforces smoothness

$$A(t) = \arg\min_{\hat{A}} \left(\left\| Y - Z\tilde{A} \right\|_2^2 + \lambda \left\| \tilde{A} \right\|_2^2 \right)$$

regularization

M>2

- Equivalent to assuming a Gaussian prior with variance determined by λ
- Large values of A are penalized. The range of allowable values for coefficients is restricted, reducing the *effective* degrees of freedom and allowing us to estimate VAR coefficients with fewer observations.

Sparsity Constraints

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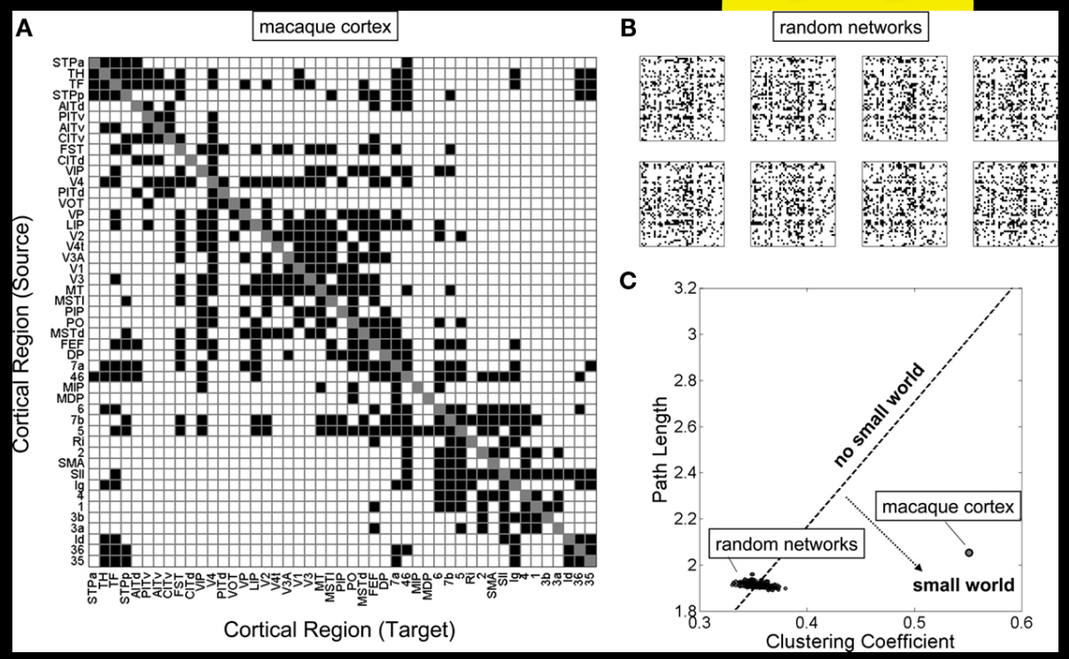
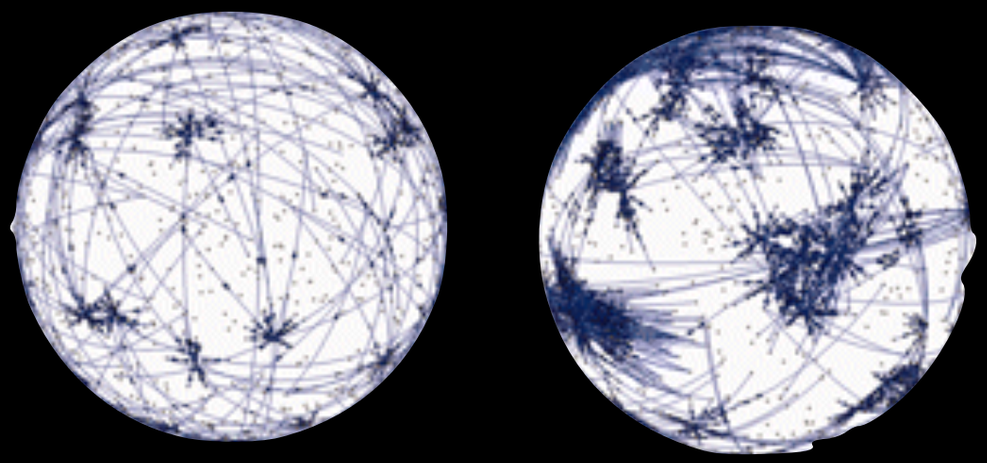
Sparsity

- Relatively low probability of a *direct* connection between any two anatomical functional units. This probability decreases with distance



Sporns, *Frontiers in Computational Neuroscience*, 2011

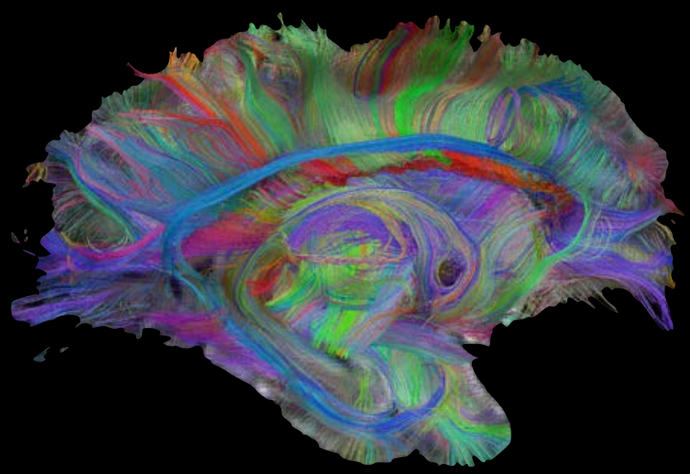
It's a small world...



structural network

functional network

Sporns and Honey, *PNAS*, 2006



Structural Connectivity

Sparsity Constraints

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- Standard least-squares solution

prediction error

$$A(t) = \arg\min_{\hat{A}} \left(\left\| Y - Z\tilde{A} \right\|_2^2 \right)$$

$$\mathbf{X}(t) = \sum_{k=1}^{p} \mathbf{A}^{(k)}(t)\mathbf{X}(t-k) + \mathbf{E}(t)$$

$$\tilde{A} = [A^{(1)}(t), \ldots, A^{(p)}(t)]^T$$

$$X_k = [X(p+1-k), \ldots, X(N-k)]^T$$

$$Z = [X_1, \ldots, X_p]$$

$$Y = X_0$$

Rewrite VAR[p] as VAR[1]

Sparsity Constraints

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- Group Lasso ($L_{1,2}$ norm)

prediction error

smoothness (L2)
(preserves spectrum)

$$A(t) = \arg\min_{\hat{A}} \left(\left\| Y - Z\tilde{A} \right\|_2^2 + \lambda \sum_{ij} \left\| \tilde{A}_{ij}^{(1)}, \ldots, \tilde{A}_{ij}^{(p)} \right\|_2 \right)$$

regularization

group sparsity (L1)

ADMM
DAL

- Equivalent to assuming a Gaussian prior over coefficients within groups and a Laplacian prior over the groups themselves
- Entire groups of coefficients are jointly pruned (set *exactly* to zero) while remaining groups assumed to have a Gaussian prior (ridge penalty). Allows us to estimate VAR coefficients with fewer observations.

Sparsity Constraints

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Compressive Sensing

- The process of acquiring and reconstructing a quantity that is underdetermined but known to be sparse (compressible) in some basis

How many samples do we need?

- N = number of samples required
- *M = number of variables/sources, p = model order*

$$N = O\left(K \log(M^2 p / K)\right) \approx O\left(\log M^2 p\right) \leq c \log\left(M^2 p\right)$$

$$N \geq M^2 p$$

(unconstrained)

Constraints Improve Estimation (if prior assumptions are correct)

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- Significant improvements using smoothness or sparsity assumptions
- (e.g. Haufe et al, 2009, Valdez-Sosa et al, 2009)

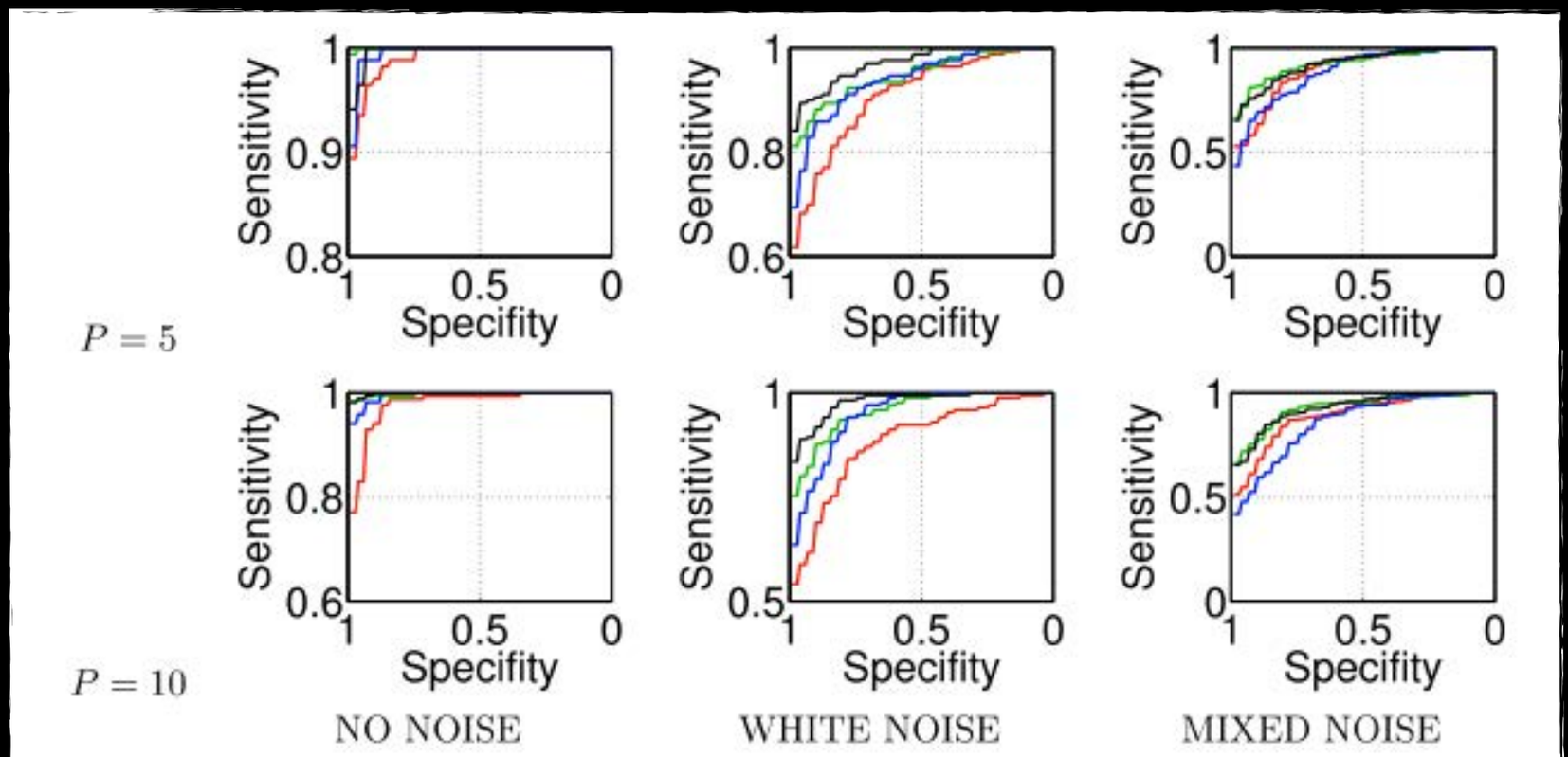

Figure 2: Average ROC curves of Granger Causality (red), Ridge Regression (green), Lasso (blue) and Group Lasso (black) in three different noise conditions and for two different model orders.

Haufe, 2009

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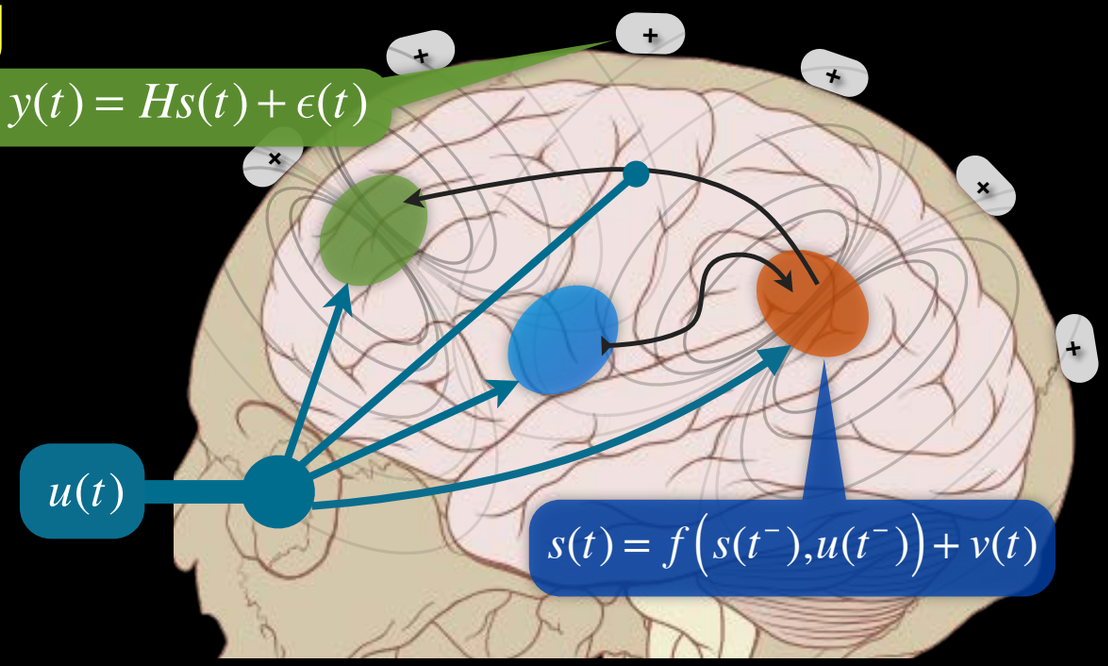
Adapting to Non-Stationarity

- **Many ways to do adaptive VAR estimation**
- Two popular approaches (adopted in SIFT):
  - Segmentation-based adaptive VAR estimation (assumes local stationarity)
  - **State-Space Modeling**

    Kalman Filtering and extensions



Discrete State-Space Model (SSM) for Electrophysiological Dynamics

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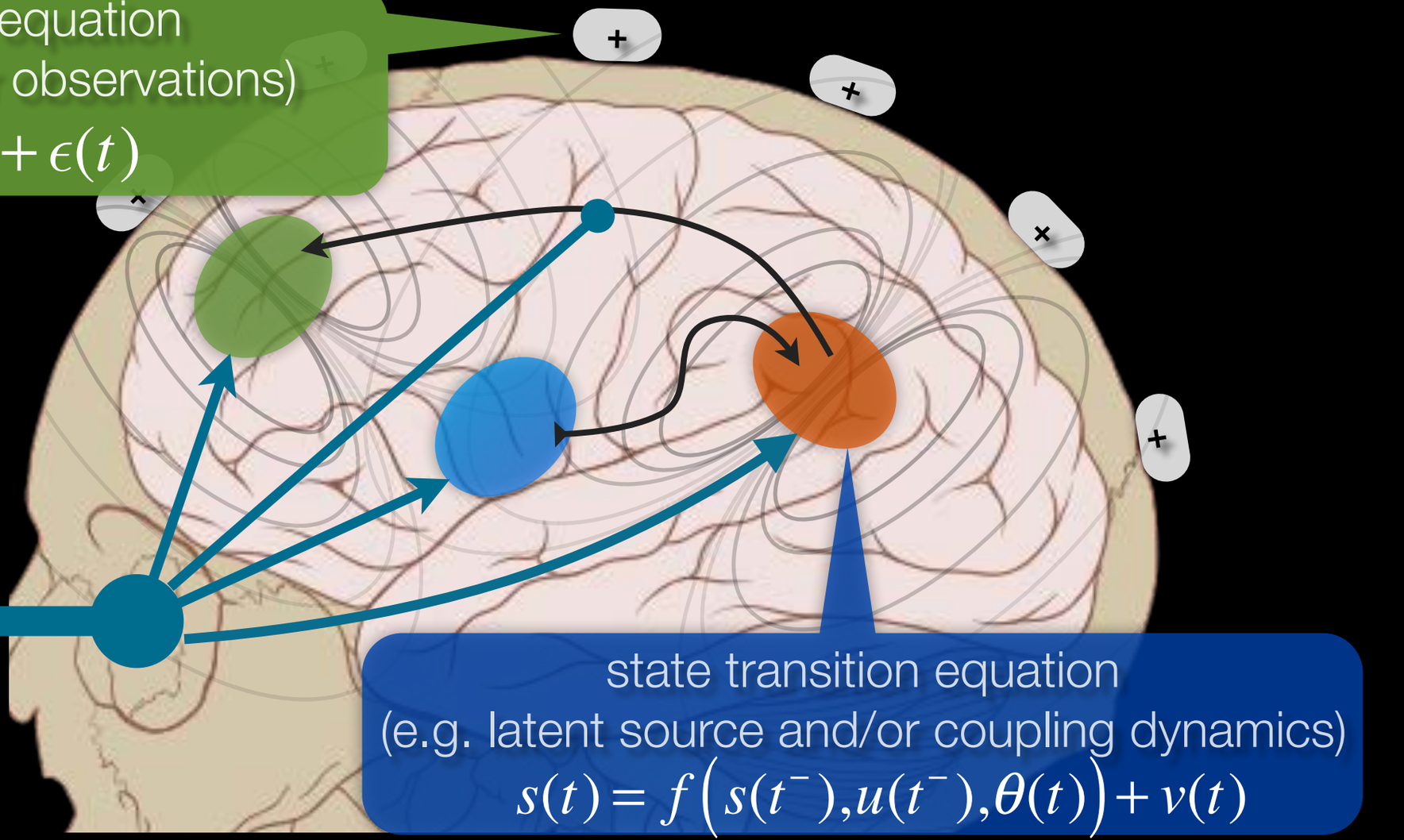
observation equation
(e.g. noisy sensor observations)

$$y(t) = Hs(t) + \epsilon(t)$$



known
deterministic inputs
$u(t)$

state transition equation
(e.g. latent source and/or coupling dynamics)

$$s(t) = f\left(s(t^-), u(t^-), \theta(t)\right) + v(t)$$

Linear VAR[1] $s(t) = \mathbf{A}(t)s(t-1) + v(t)$

- Dynamical system may be linear or nonlinear, dense or sparse, non-stationary, high-dimensional, partially-observed, and stochastic
- Subsumes discrete Delay Differential Equation (DDE) and Vector Autoregressive (VAR) methods and closely related to Dynamic Causal Modeling (DCM)

Kalman Filtering

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optimal estimator (in terms of minimum variance) for the state of a linear dynamical system

$f(z,u)$ is linear in $\{z,u\}$ for classic Kalman Filtering

$\hat{y}(t)$

new observation

$y_t \quad u_t$

Time Update ("Predict")

$$\hat{z}_t^- = f(\hat{z}_{t-1}, u_{t-1}) + v_t$$
$$\hat{y}_t = H\hat{z}_t^-$$
$$P^-_t = AP_{t-1}A^T + Q$$

Measurement Update ("Correct")

$$\epsilon = y_t - \hat{y}_t$$
$$\hat{z}_t = \hat{z}_t^- + G_t\epsilon$$
$$P_t = (I - G_tH)P^-_t$$

$t \rightarrow t+1$

Initialize

$$\hat{z}_0 = E(z_0)$$
$$P_0 = \mathrm{cov}(z_0)$$

updated model

z_t := unknown state vector at time t
e.g. delay-embedding of sources and/or coupling (VAR) parameters

Kalman Filtering

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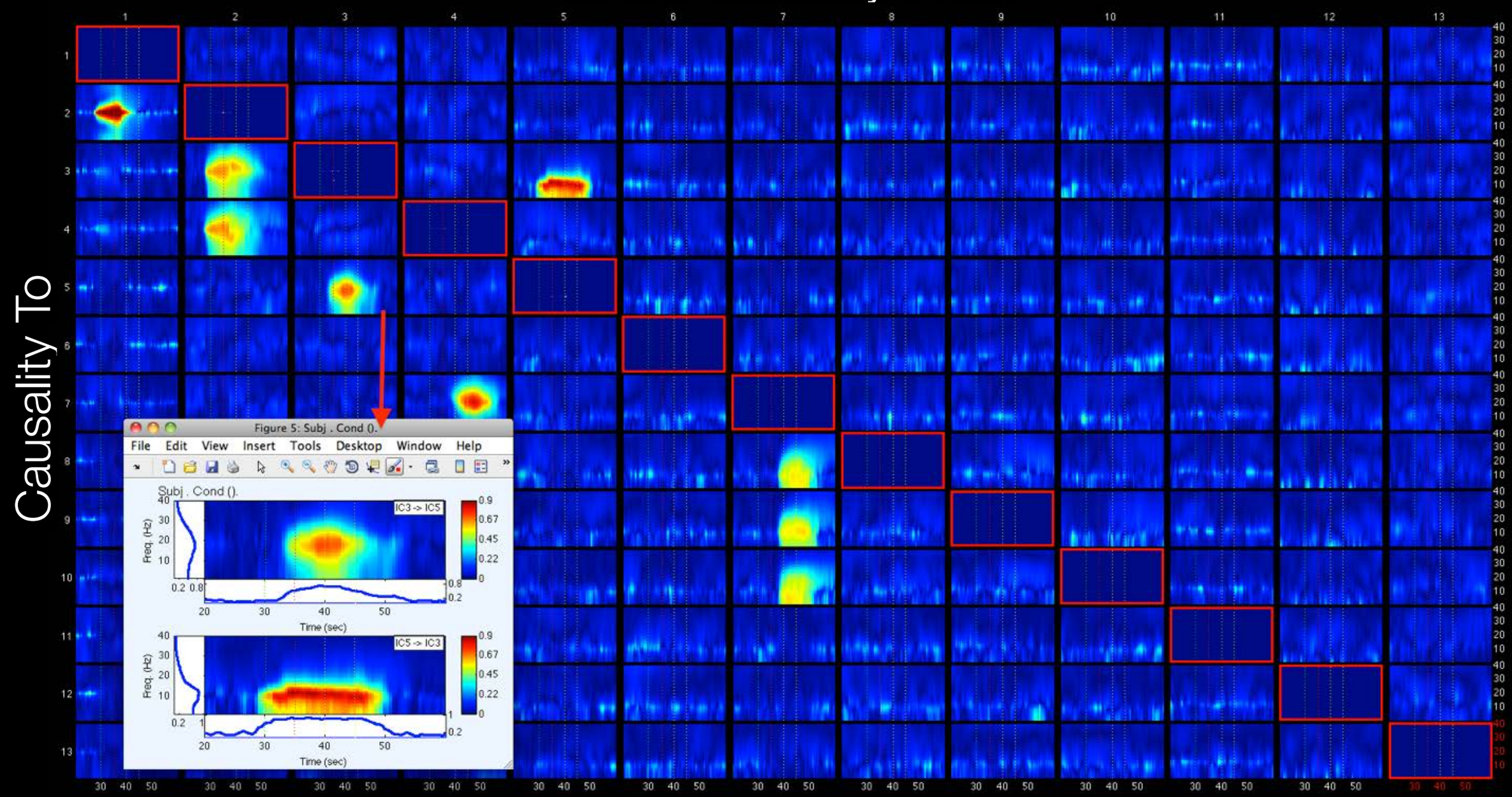

Nonlinear Modeling

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- Interactions in brain are generally non-linear
- Purely linear models (e.g. high-order VAR models) can sometimes provide an approximation sufficient for correct detection of directed dependencies

Schelter B, Timmer J, Eichler M., *J. Neuro. Methods* 2009

van-der-Pol coupled oscillators

$$\ddot{x}_i = \mu(1 - x_i^2)\dot{x}_i + \omega_i^2 x_i + \sigma_i \eta_i + \sum_{j \neq i} \epsilon_{ij}(x_j - x_i)$$

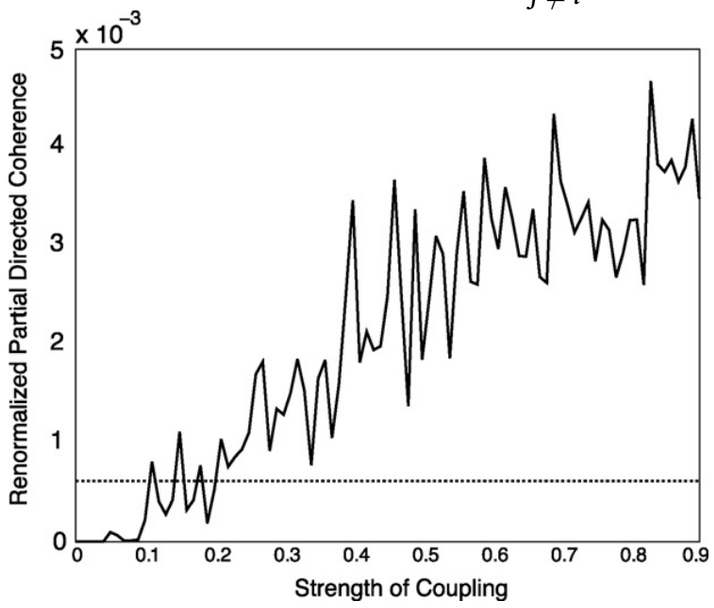

Fig. 8. Coupled van-der-Pol oscillators. Renormalized partial directed coherence for various coupling strengths ϵ_{12}. The dashed horizontal line marks the 5% significance level.

Stochastic coupled Rössler oscillators

$$\dot{\xi}_j = \begin{pmatrix} \dot{X}_j \\ \dot{Y}_j \\ \dot{Z}_j \end{pmatrix} = \begin{pmatrix} -\omega_j Y_j - Z_j + \left[\sum_{i, i \neq j} \varepsilon_{j,i} \left(X_i - X_j \right) \right] + \sigma_j \eta_j \\ \omega_j X_j + a Y_j \\ b + (X_j - c) Z_j \end{pmatrix}$$

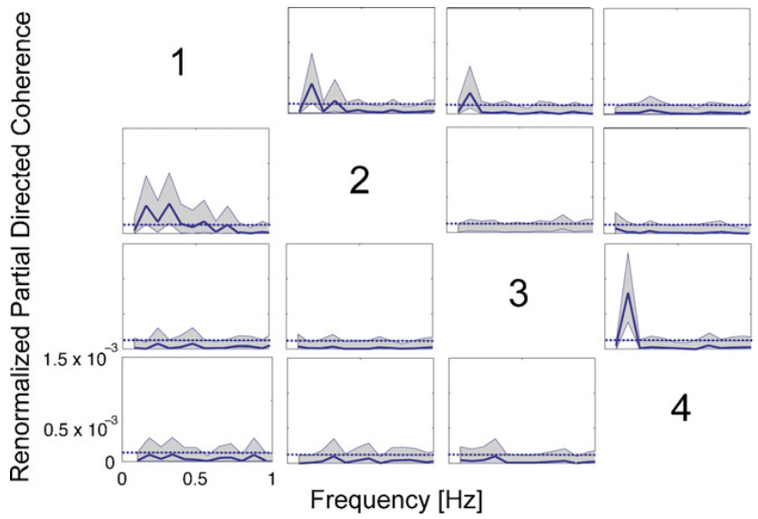

Fig. 9. Results of the renormalized partial directed coherence analysis for a network coupled Rössler oscillators. The coupling is bidirectional between oscillators 1 and 2 and unidirectional from oscillator 2 to oscillator 4 and from oscillator 4 to oscillator 3. The dashed horizontal line marks the 5% significance level, while the gray area represents the 95% confidence intervals.

Nonlinear Modeling

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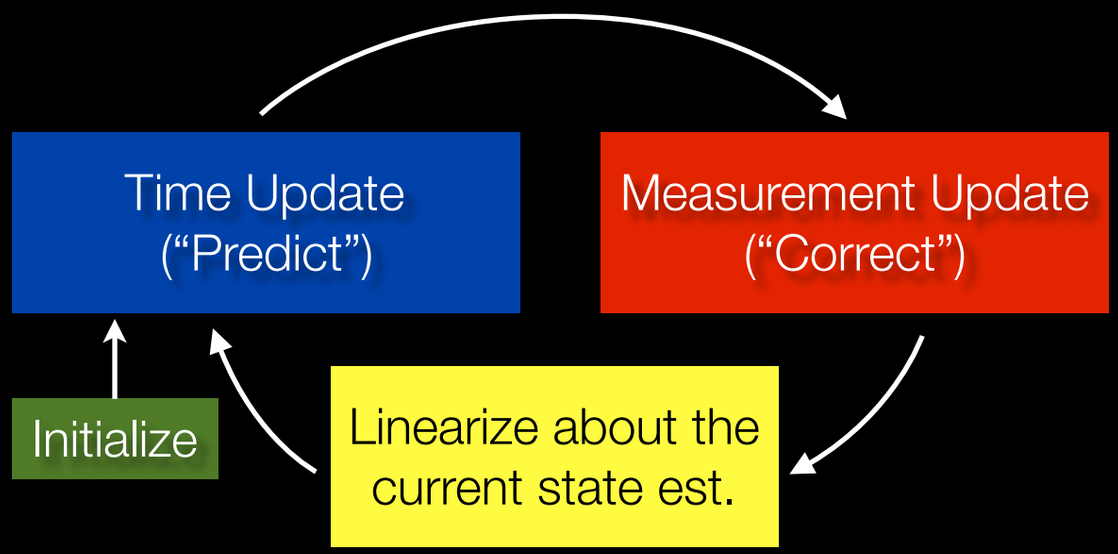
- A more general approach is to transform the non-linear system to a linear representation and apply the linear model there.
- e.g. **local-linearization**, kernel methods, etc.
- Nonlinear extensions of Kalman filtering provide efficient ways to model the time-evolving states and parameters of nonlinear processes
- (Dual) Extended Kalman Filtering



- Linearize about the current state (first-order Taylor approximation).
- Apply the Kalman Filter update rules using the linearized model
- Apply Granger-Geweke Causality to the linearized coefficient matrices

Kalman Filtering in SIFT

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- Linear Kalman Filter: est_fitMVARKalman.m
- Nonlinear Kalman Filter: est_fitMVAR_DEKF.m

Statistics

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- Different ways to do statistics in SIFT
  - Phase Randomization
  - Bootstrapping
  - Analytic Tests

Test	Null Hypothesis	What question are we addressing?	Applicable Methods
H_{null}	$C(i,j) = 0$	Is there significantly non-zero information flow from process $j \longrightarrow i$?	Phase randomization Analytic tests
H_{base}	$C(i,j) = C_{base}(i,j)$	Is there a difference in information flow relative to the baseline?	Bootstrap resampling
H_{AB}	$C_A(i,j) = C_B(i,j)$	Is there a difference in information flow between experimental conditions/populations A and B?	Bootstrap resampling

C(i,j) is the measured information flow from process j --> i.
C_{null} is the expected measured information flow when there is no true information flow.
C_{base} is the expected information flow in some baseline period.

Statistics

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Statistical Approach	Test	Parametric	Nonparam.
Asymptotic analytic estimates of confidence intervals. Applies to: PDC, nPDC, DTF, nDTF, rPDC	H_{null}, H_{base}, H_{AB}	☑	
Theiler phase randomization Applies to: all	H_{null}		☑
Bootstrap, Jacknife, Cross-Validation Applies to: all	H_{AB}, H_{base}		☑
Confidence intervals using Bayesian smoothing splines Applies to: all	H_{base}, H_{AB}	☑	☑

$H_{null} : \mathbf{C}_{ij} = 0$ $H_{base}: \mathbf{C}_{ij} = \mathbf{C}_{baseline}$ $H_{AB}: \mathbf{C}^{A}_{ij} = \mathbf{C}^{B}_{ij}$

Bootstrap Statistics

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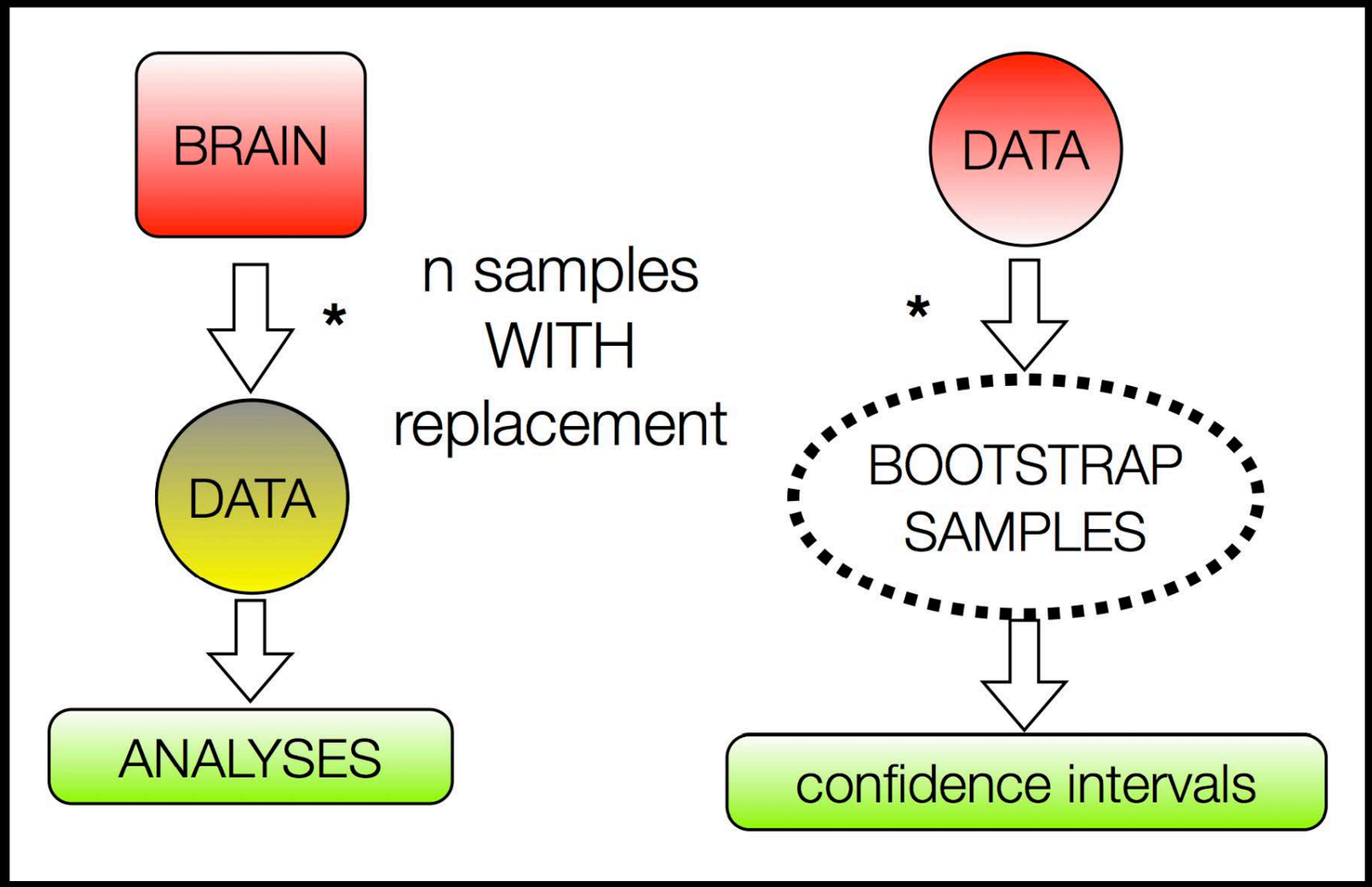

- sample = X1, ..., Xn
- for k=1:R (number of bootstrap resamples/iterations)
  - resample n observations (trials) with replacement $X^* = \{X^*_{1, \ldots} X^*_n\}$
  - compute estimator E_k (fit model, obtain connectivity) based on X^*
  - repeat
- with R large enough $P_E = \{E_1, \ldots, E_R\}$ provides a good approximation to the true distribution of the estimator (connectivity, power, etc)

Bootstrap Statistics

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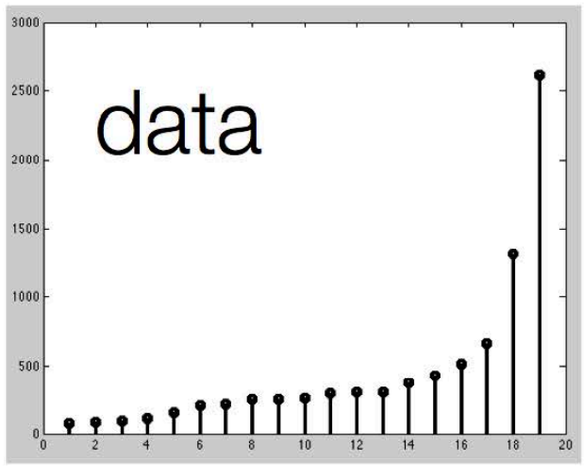
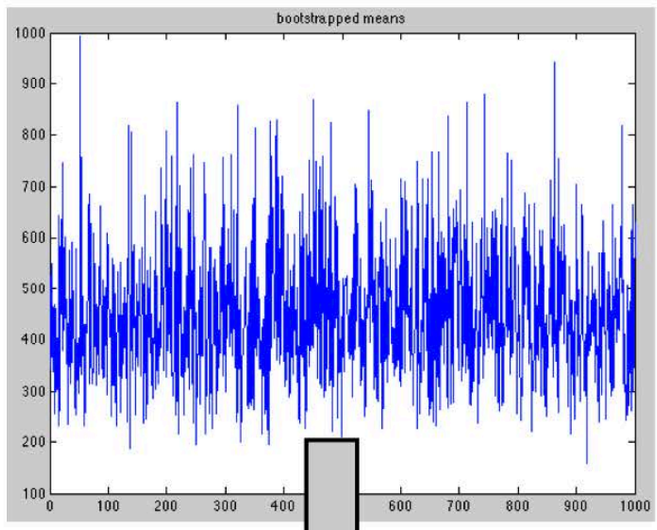
% self-awarness data, Wilcox, 2005, p58



Sample with replacement b times

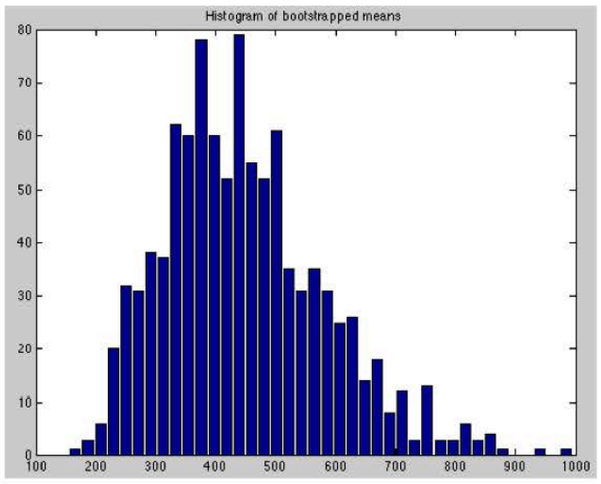
Bootstrapped estimates



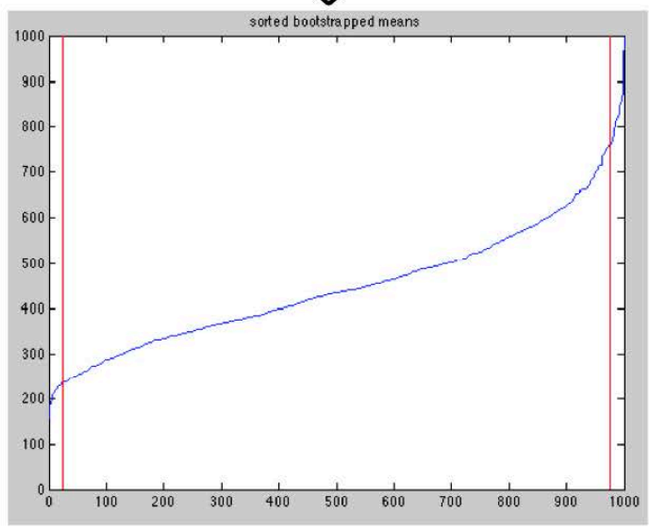
compute estimate

get PDF

Distribution of bootstrapped estimates of the mean



Sort & get CI



Bootstrap Statistics

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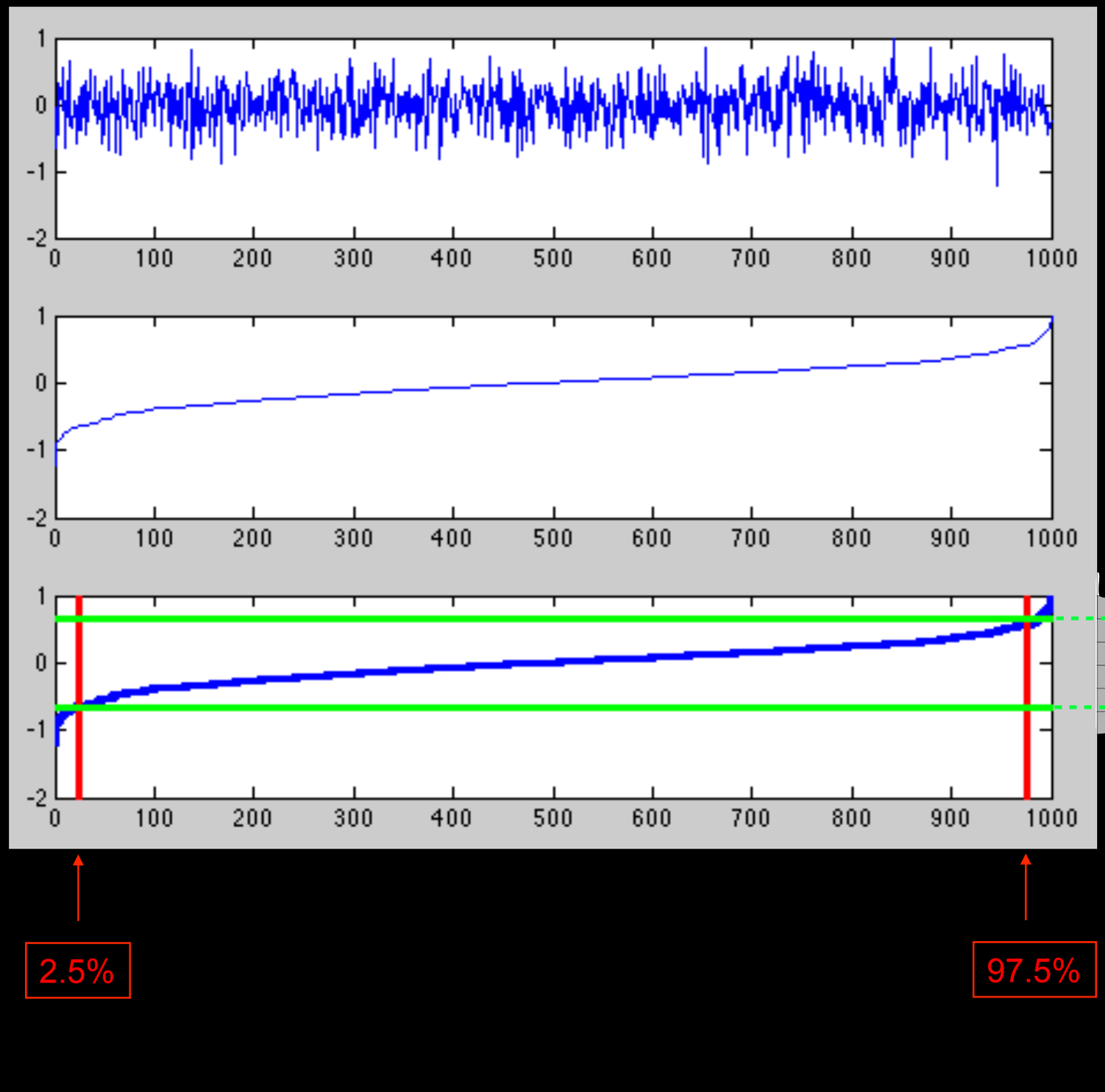
bootstrap

sorted values (cdf)

thresholds (ci)



Bootstrap Differences

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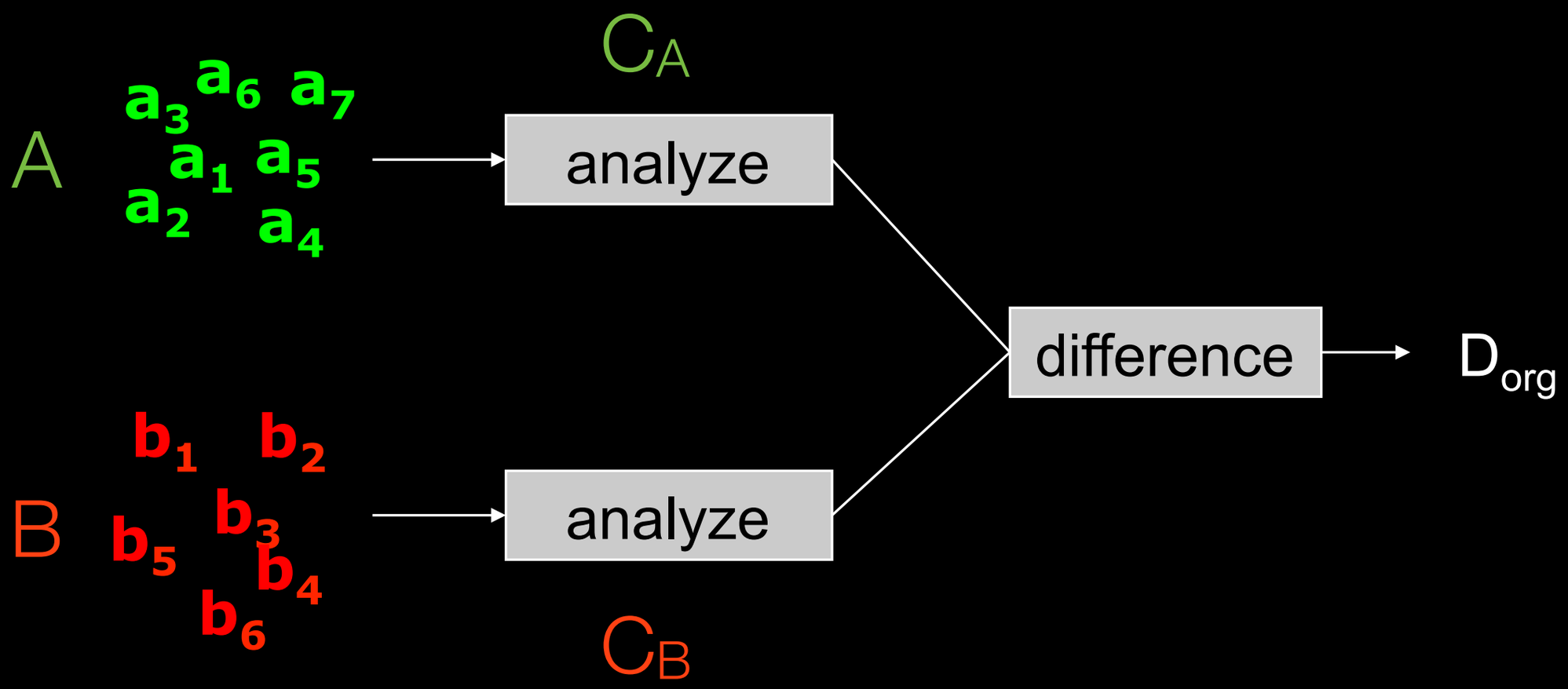
- Suppose we have two conditions
  A = {a1,...a7}
- B = {b1,...,b6}
- We want to estimate the distributions of connectivity estimator applied to A and B separately, as well as the difference distribution (for testing H0: A=B)



Bootstrap Differences

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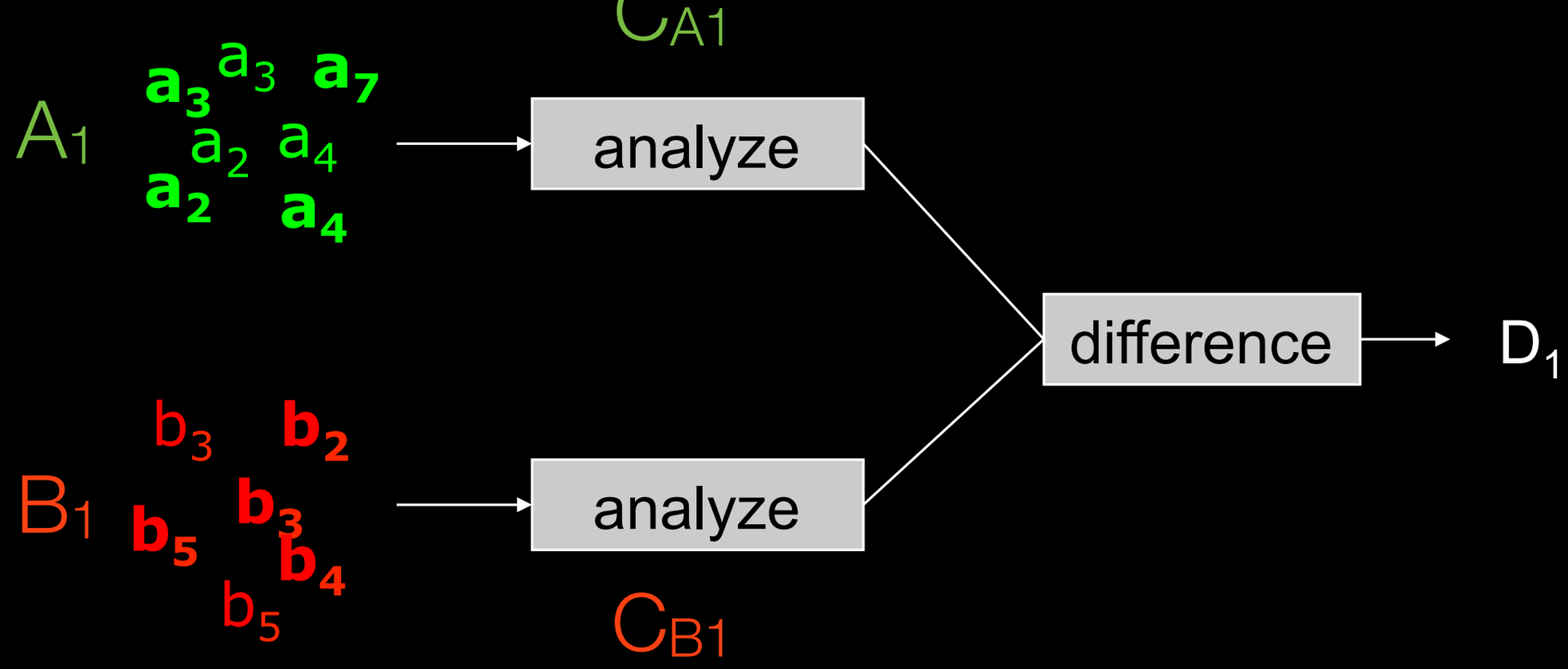
- For k=1:R (number of bootstrap iterations)
  - Resample with replacement from both groups to get A_k and B_k
  - Fit models and obtain connectivity C_{Ak}, C_{Bk}
  - Compute difference $D_k = C_{Ak}-C_{Bk}$
- Repeat



Bootstrap Differences

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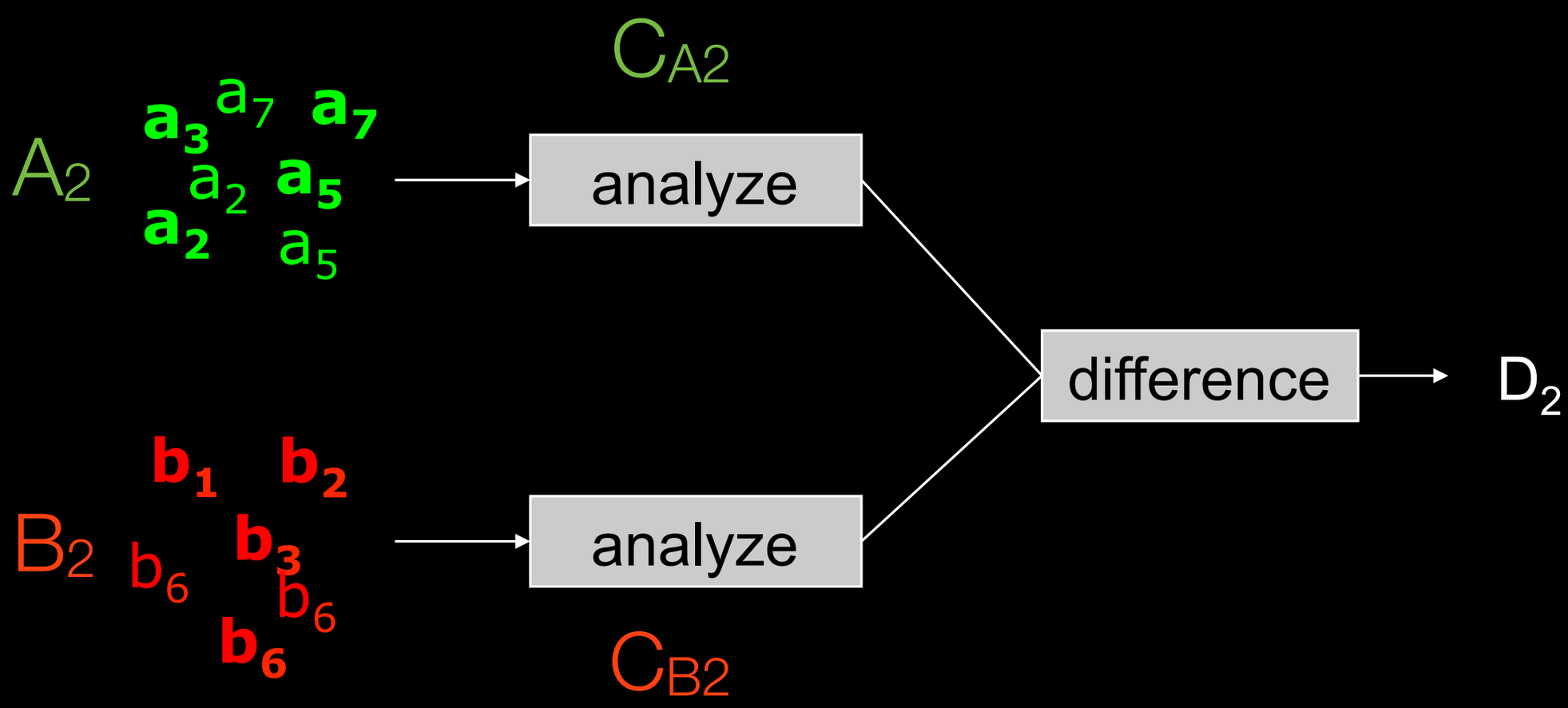
- For k=1:R (number of bootstrap iterations)
  - Resample with replacement from both groups to get A_k and B_k
  - Fit models and obtain connectivity C_{Ak}, C_{Bk}
  - Compute difference $D_k = C_{Ak}-C_{Bk}$
- Repeat



Bootstrap Statistics

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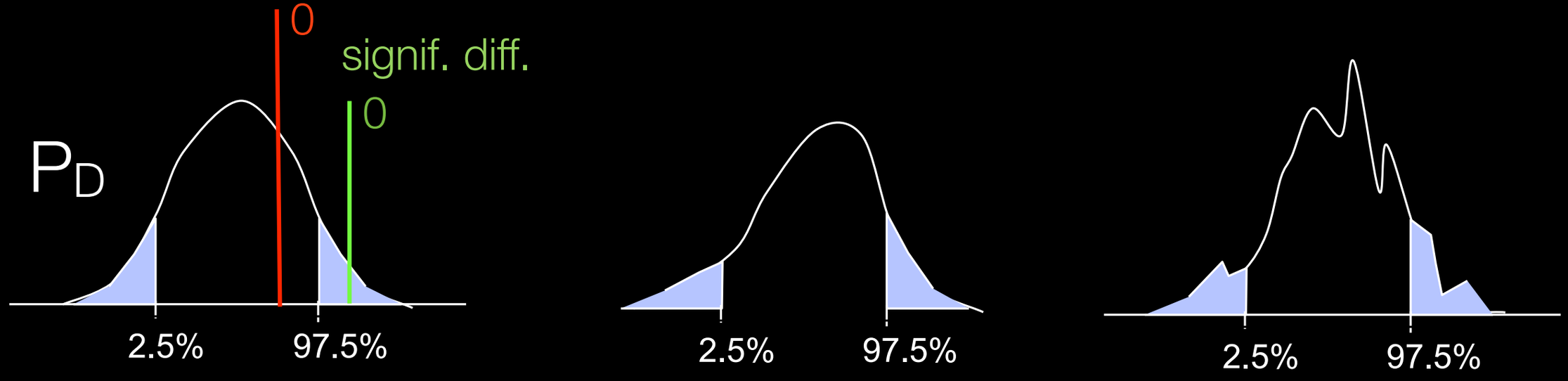
- The procedure yields a distribution $P_D = \{D_1, ... D_R\}$
- If 0 lies in the right (or left) tail of this "difference distribution", then we **reject** the null hypothesis that A=B at the chosen confidence level (below: alpha=0.05 for a two-sided test)



- Difference distribution can take any shape
- The procedure above also provides estimates of the individual distributions of C_A and C_B yielding confidence intervals for H1

Phase-Randomization

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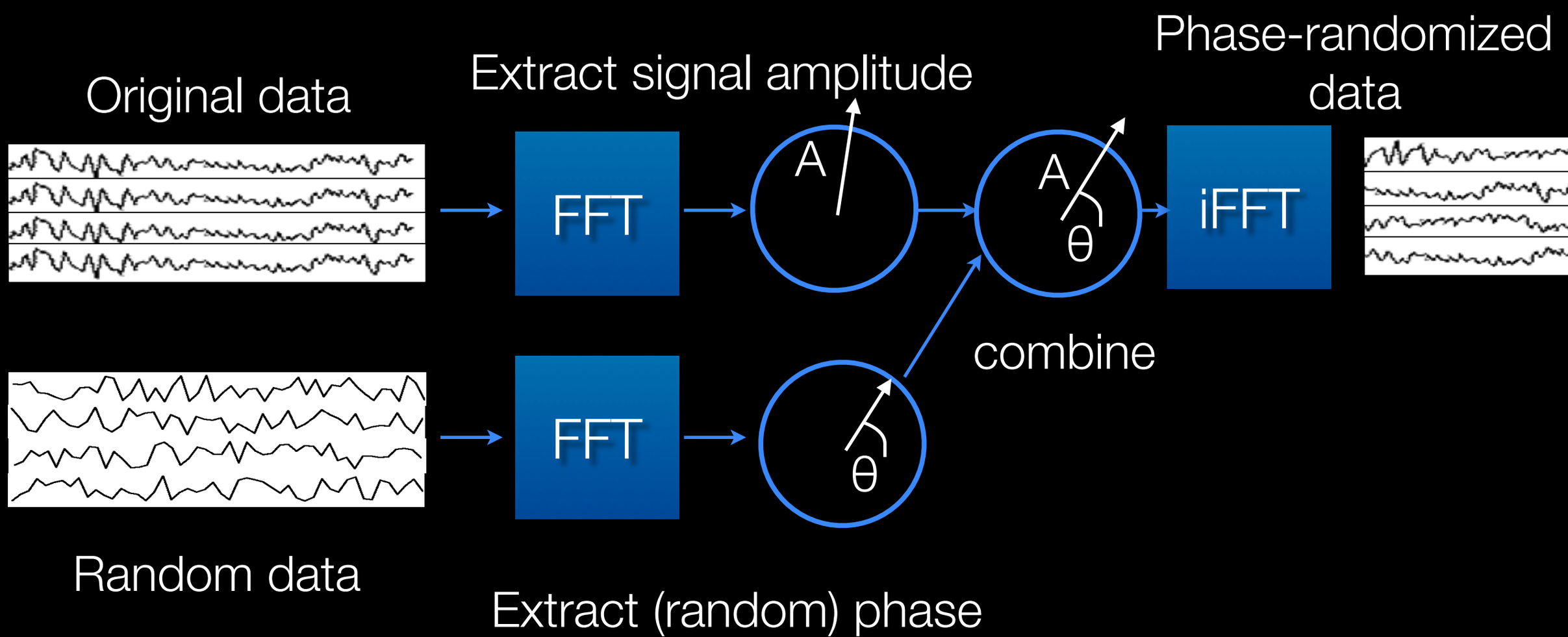
- Phase Randomization Procedure (Theiler, 1992)
  - Method for testing whether there is non-zero information flow (H_{null})



Phase-Randomization

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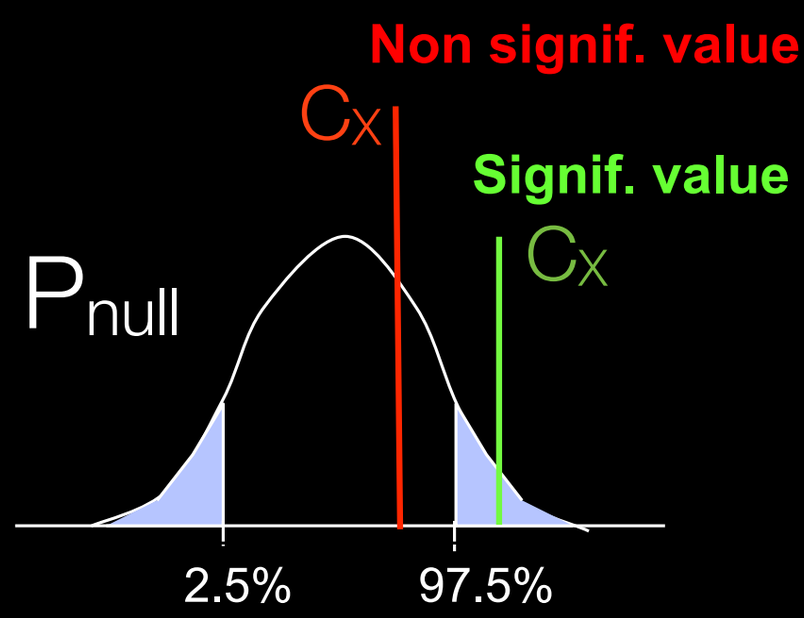
- Start with an n-trial sample: X = {X1, ..., Xn}
- for k=1:R (number of resamples)
  - randomize phases for all trials
  - compute connectivity estimate C_k
  - repeat
- With B large enough the B estimates provide a good approximation of the **null** distribution of the connectivity estimator
- Compare connectivity C_X from original (non-randomized) samples X to quantiles of $P_{null} = \{C_1, ..., C_R\}$



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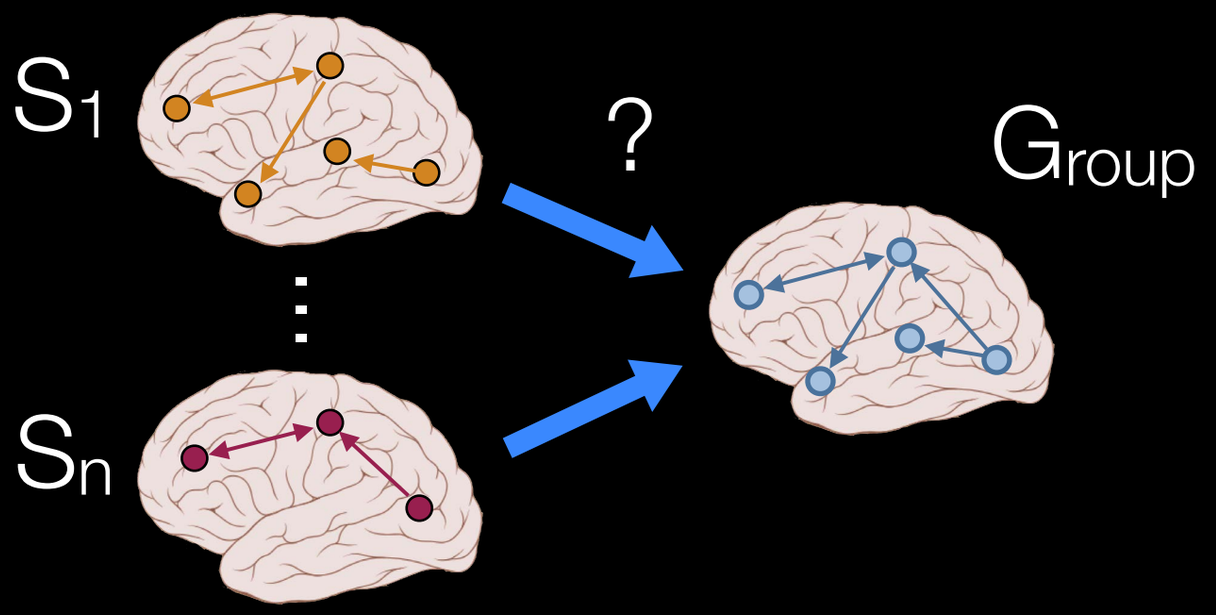
Multi-Subject Inference

- In many cases of source analysis involving focal/point sources (e.g. BSS + dipole fitting, sparse patch-based estimation) we encounter two key problems w.r.t. multi-subject inference:

  1. Identification/Co-registration
  2. Missing Data



- Conventional approaches utilizing disjoint clustering do not suitably address Issue 2 and generally lack means for rigorously quantifying statistical uncertainty in addressing Issue 1

  - Issue 2 is exacerbated in connectivity analysis due to the combinatorial explosion of variables associated with a given source

Bayesian Hierarchical Model

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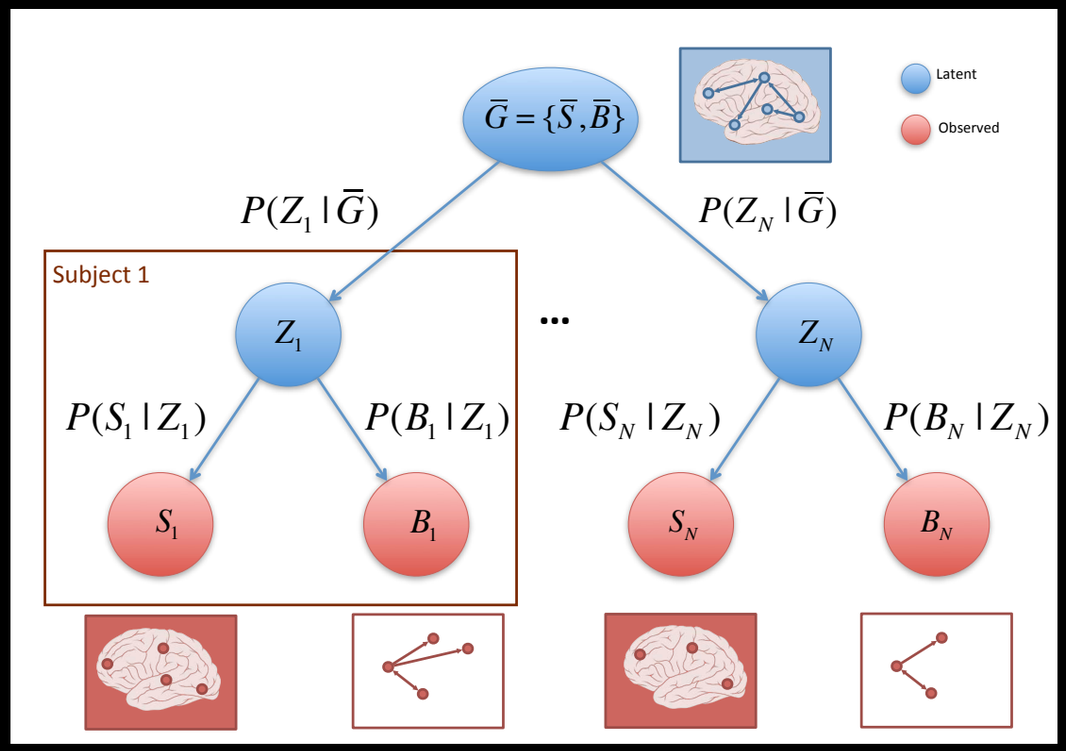
- Perform multi-subject (second-level) inference via hierarchical (mixture) model, approximating the posterior distribution of source locations and connectivity surfaces.

- Advantages:
  - Handles multivariate measures such as connectivity
  - Yields posterior distributions allowing robust statistics and increased range of hypothesis testing (incl. analysis of individual variability)
  - Hierarchical structure can be adapted to obtain conditional probability distribution w.r.t. other metrics (ERSP, genetics, morphometry, behavior, etc)



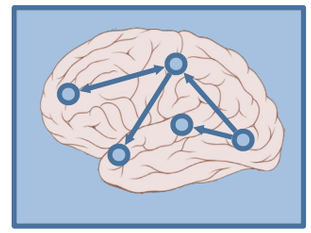
See: Mullen, T. "The Dynamic Brain: Modeling Neural Dynamics and Interactions From Human Electrophysiological Recordings" Chapter 3.

Generative Model

Goal:
$P(G,Z \mid \{S,B\})$

Group-Level Network

$\bar{G} = \{\bar{S}, \bar{B}\}$



Latent

Observed

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$P(Z_1 \mid \bar{G})$ $P(Z_N \mid \bar{G})$

Indicator variables assigning subject-level sources to group-level clusters

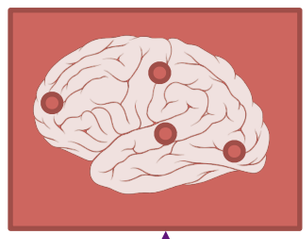
Subject 1

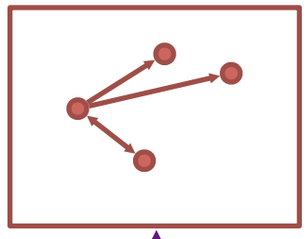
Z_1 ... Z_N

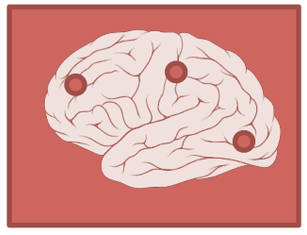
$P(S_1 \mid Z_1)$ $P(B_1 \mid Z_1)$ $P(S_N \mid Z_N)$ $P(B_N \mid Z_N)$

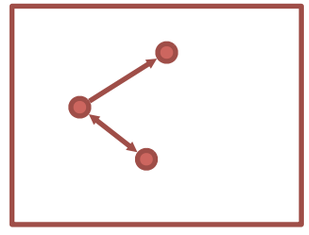
Observed Data

S_1 B_1 S_N B_N

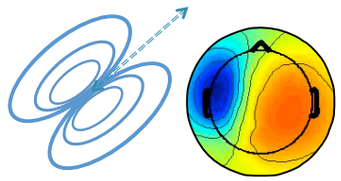








$S \in \{x,y,z\}$ $B = E\{\mathbf{b}\}$



Source Localization

Source Separation

$\mathbf{Y}(t)$

B-spline

Conn

MVAR

Bayesian FPCA Smoothing (Parameterization)

$$\mathbf{C}(f,t) = g(f,t,\mathbf{b}) + \eta(f,t)$$

$$\mathbf{C}(f,t) = q(\mathbf{A}^{(k)}(t), \mathbf{E}(t))$$

$$\mathbf{Y}(t) = \sum_{k=1}^{p} \mathbf{A}^{(k)}(t)\mathbf{Y}(t-k) + \mathbf{E}(t)$$

MCMC Estimation (Gibbs Sampling)

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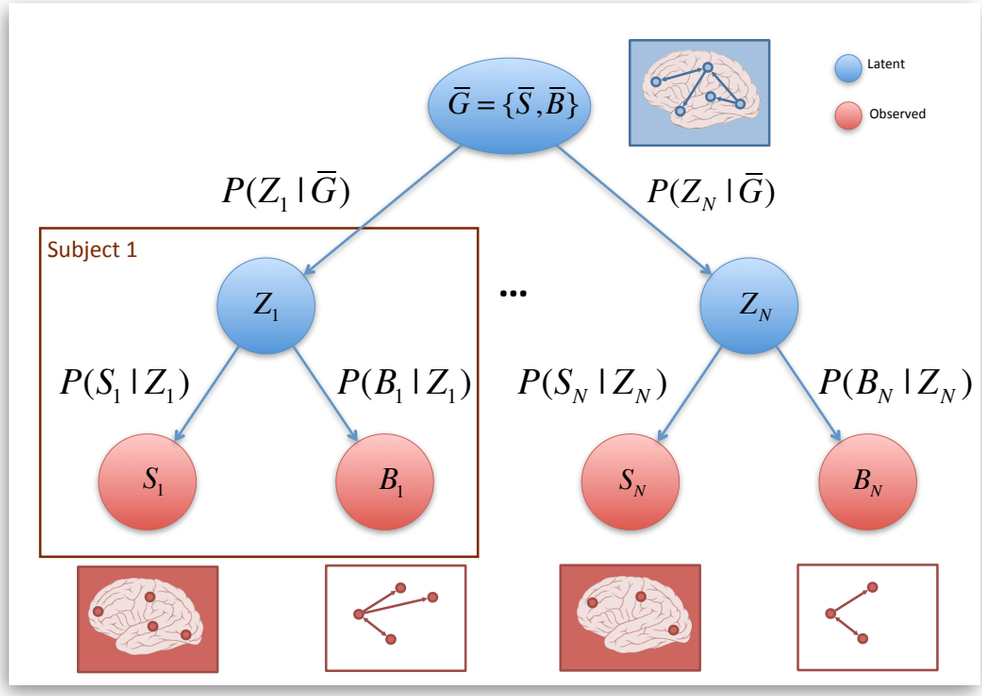
$$D = \{\{S_i\},\{B_i\},\{\Sigma_s\},\{\sigma_b{}^2\}\}$$

$$G = \{\bar{S},\bar{B},\Sigma_{\bar{s}},\sigma_{\bar{b}}{}^2\}$$



$P(G,Z \mid D)$ Target Distribution

$$P(S_i \mid Z_i, G) = \prod_{j=1}^{M_i}\prod_{k=1}^{M}\left[N(s_{ij} \mid \bar{s}_k, \Sigma_{s,k})\right]^{z_{ijk}}$$

$$P(B_i \mid Z_i, G) = \prod_{j_1=1}^{M_i}\prod_{j_2=1}^{M_i}\prod_{k_1=1}^{M}\prod_{k_2=1}^{M}\left[N(b_{ij_1j_2} \mid \bar{b}_{k_1k_2}, \sigma^2_{\bar{b},k_1k_2} I)\right]^{z_{ij_1k_1} z_{ij_2k_2}}$$

Gibbs Sampling Scheme:

1) Initialize G^*, Z^* from initial clustering (e.g. k-means) solution

2) Sample G^* from posterior $P(G^* \mid Z^*, D) \propto P(D \mid Z^*, G^*)P(G^*)$

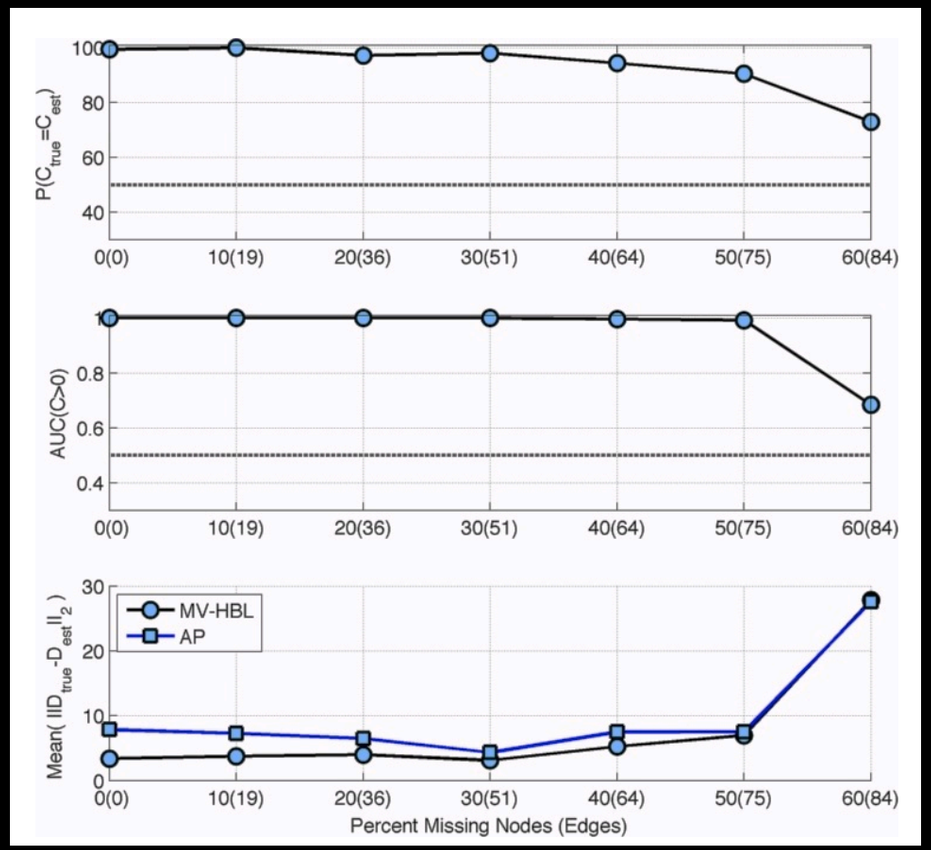
3) Sample Z^* from posterior $P(Z^* \mid G^*, D) \propto P(D \mid Z^*, G^*)P(Z^*)$

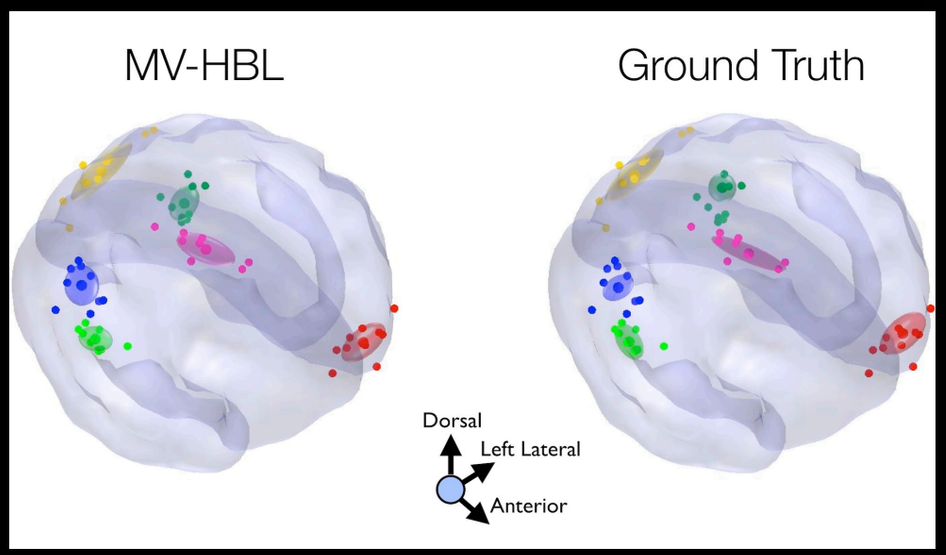
4) Repeat (2-3) many times: $P(G^*,Z^* \mid D) \rightarrow P(G,Z \mid D)$

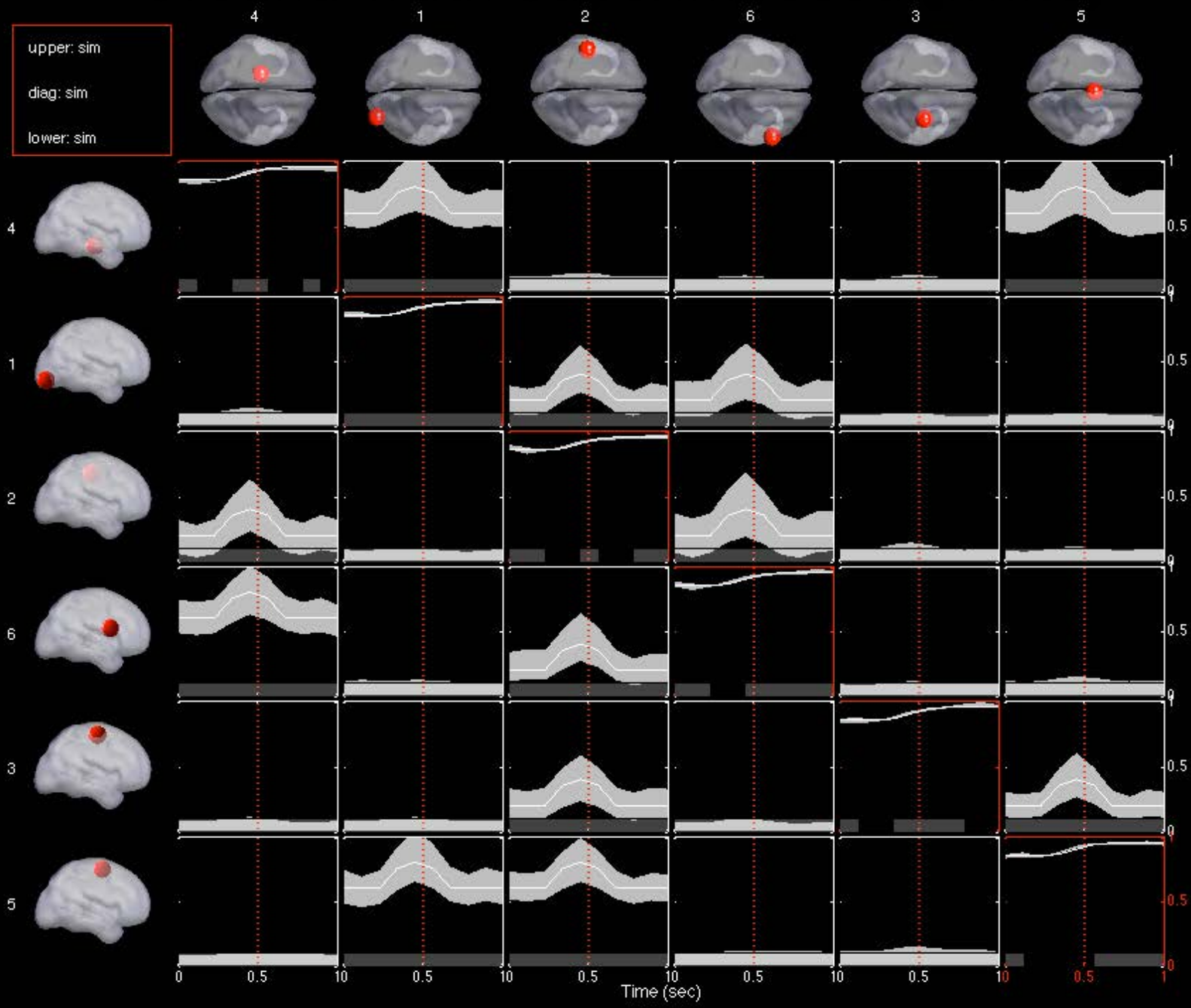
MV-HBL

Sim: 10 Subjects, 30% missing dipoles

Estimated 95% CI (gray) with Ground Truth Superimposed (white)







Est = True ($p < 0.05$)

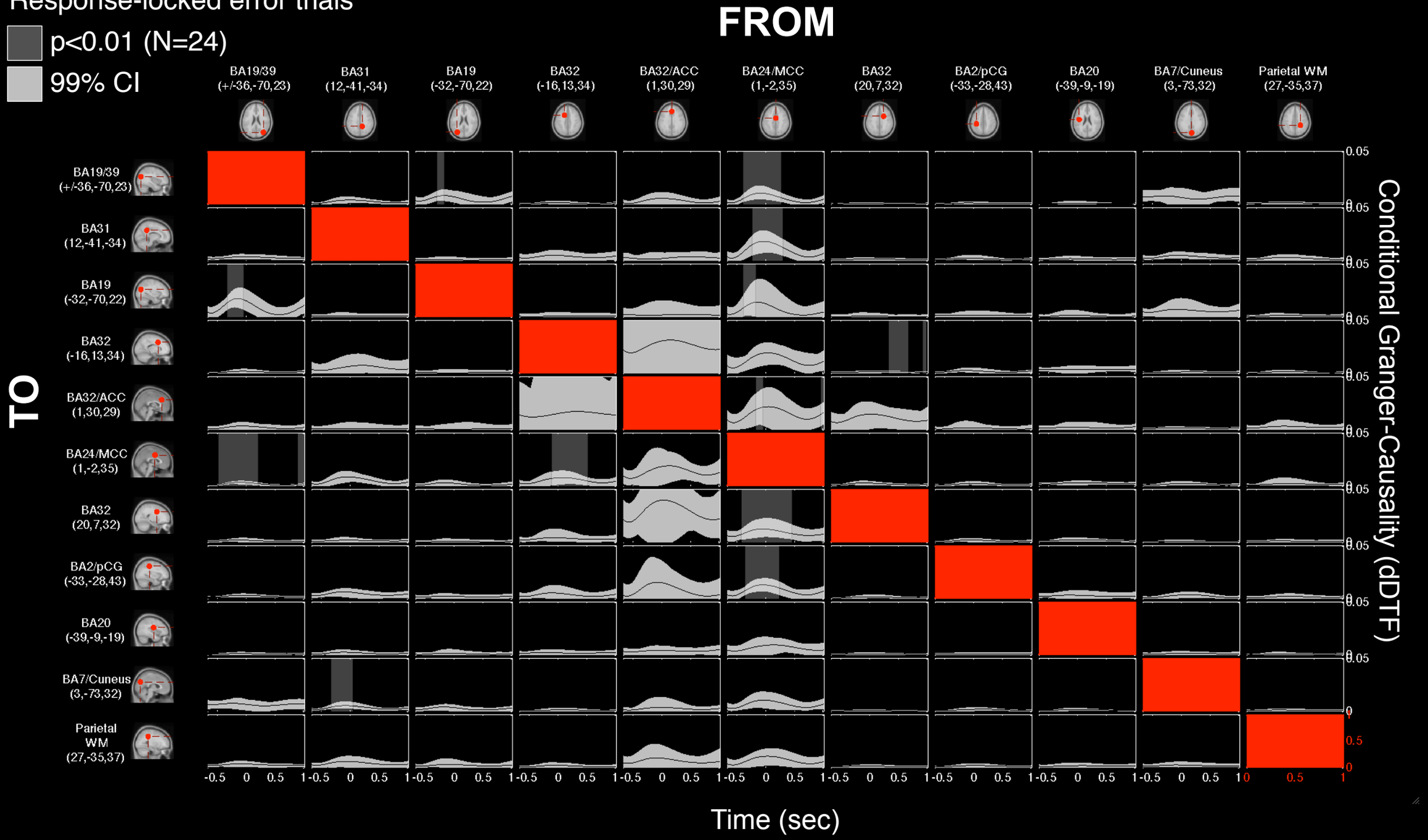
Est. 95% CI

Mullen, T., 2014 “The Dynamic Brain”; Mullen & Thompson, in prep

Bayesian Multi-Subject Inference

Swartz Center for Computational Neuroscience
Tim Mullen

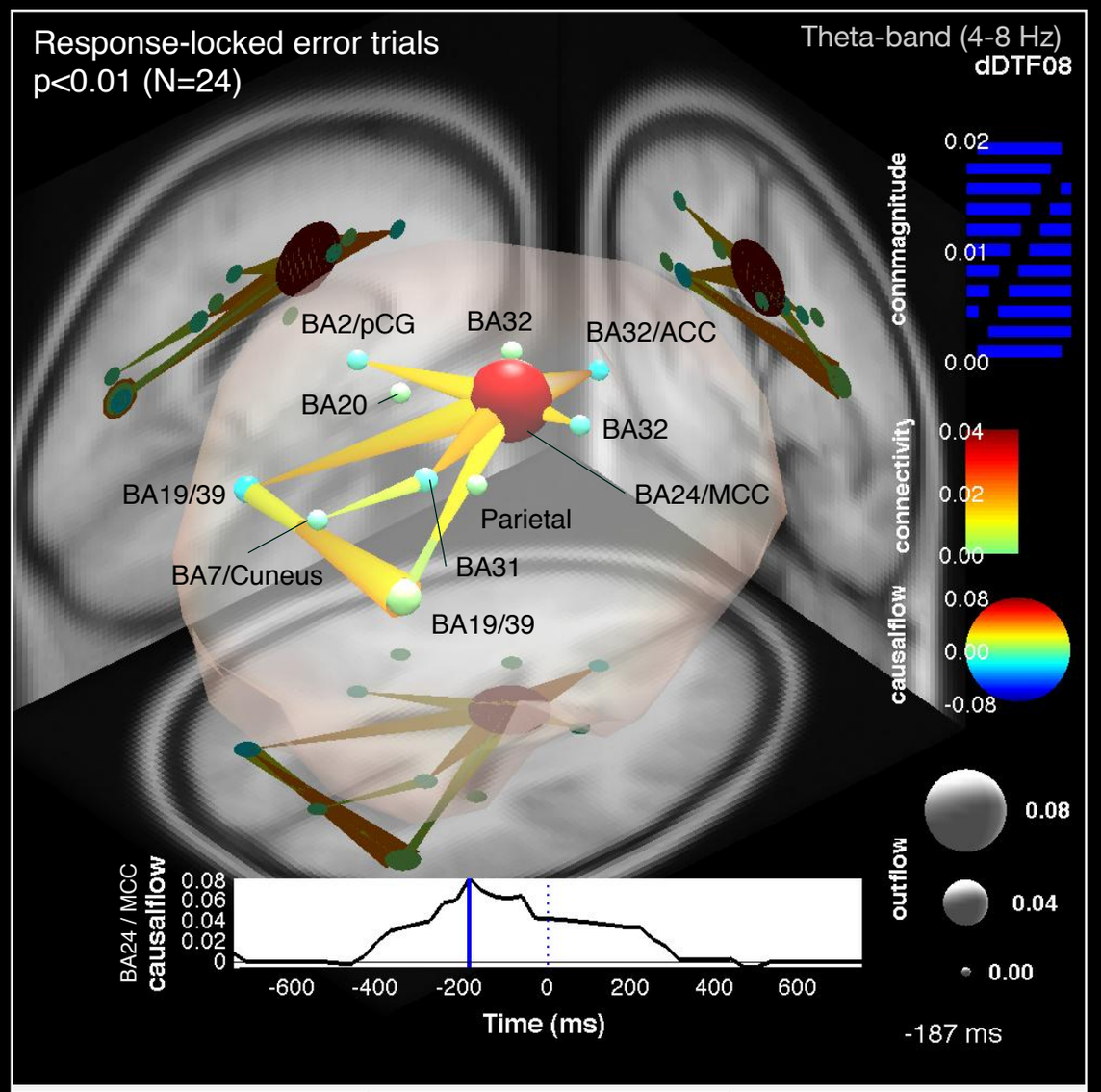
Theta-band (4-8 Hz) event-related dDTF
Baseline: [-750 -500] ms
Response-locked error trials
p<0.01 (N=24)
99% CI

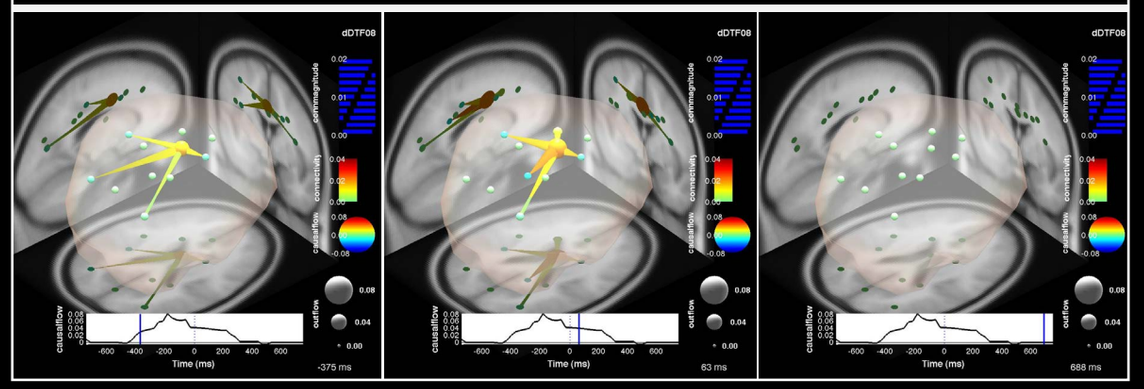


Cluster retained if more than 33% of subjects have greater than 50% probability of cluster membership

Bayesian Multi-Subject Inference

Swartz Center for Computational Neuroscience
Tim Mullen





Cluster retained if more than 33% of subjects have greater than 50% probability of cluster membership

Conventional Statistics

Swartz Center for Computational Neuroscience
Tim Mullen

- An alternative approach:
  For each subject…

  1. Perform distributed source localization (possibly after separating a subspace of brain components using ICA)

  2. Select *M* regions of interest (ROIs) e.g. from a standardized anatomical atlas (e.g. Desikan-Killiany, Destrieux, etc) and integrate current density within each ROI. This yields *M* source time-series for each subject

  3. Store results in EEG.srcpot

  4. Obtain connectivity estimates for sources using SIFT with the 'Sources' option set in pre-processing. Resulting [M x M x N_{freq} x N_{times}] connectivity matrices are stored in EEG.CAT.Conn.

5. Apply your favorite mass-univariate or multivariate statistical approach (e.g. GLM, t-test, (M)ANOVA, etc) to the collection of connectivity estimates from all subjects to obtain desired statistics. See LIMO-EEG Toolbox and EEGLAB's statcond(). Beware of multiple comparisons issues! FDR may not be suitable.