

## Clustering Independent Components of EEG Data



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# Why cluster independent components across subjects or sessions?



- ICA transforms the data from a channel basis
  (activity recorded at each channel)
  - to a component basis (activity computed at each IC).
- Normally, EEG researchers assume that, for example, electrode channel F7 == F7 == F7 ... in each subject – and then 'cluster' their data assuming channel equivalence.
- This amounts to the simple assumption

"Your Cz is My Cz!"

But this is only *roughly* correct !













## Machine Clustering ICA components by eye





Makeig et al., ~2000 unpublished

#### Man So how to cluster components?



#### The same problems hold for clustering independent components

Across Ss, components don't even have "the same" scalp maps! → Are "the same" components found across subjects?

- What should define "*the same*" (i.e., "*component equivalence*")?
  - Similar scalp maps?
  - Similar cortical or 3-D equivalent dipole locations?
  - Similar activity power spectra?
  - Similar ERPs?

- Similar ERSPs?
- Similar ITCs?
- Or similar *combinations* of the above?? ...

# EEGIC Source Locations



#### (135,794 IC equivalent dipoles!)



#### ... Some caveats

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#### In this *dipoledensity()* assay ...

- MR head images were not available  $\rightarrow$  brain co-registration crude.
- Single versus dual-dipole model selection was subjective.
- Different electrode montages  $\rightarrow$  mis-localization effects.
- Electrode locations were not all digitized some 'guestimated' !
- Brain geometries differ!





### Co-Registration of Electrodes with MR Image







**EEG** 

#### MR + EEG

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Onton & Makeig, 2004





## Does the spatial distribution of IC equivalent dipole source locations depend on the task the subject performs?

i.e.

## Do "the same" ICs (and IC clusters) appear for every task?

## www.www.**Equivalent dipole density**





>> dipoledensity()

Onton et al., 2

Onton et al., '05

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>> dipoledensity()

Onton et al., 2

Onton et al., '05

## www.www.**Equivalent dipole density**





Sternberg letter memory task

>> dipoledensity()

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Emotion imagery task

>> dipoledensity()

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  - Similar ERSPs?
  - Similar ITCs?
  - Or similar *combinations* of the above?? ...
  - EEGLAB clustering supports all these possibilities.



## **Study IC Clustering: Assumptions**

• Assumes there are *functionally equivalent* ICs across most subjects.

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- Assumes these ICs have similar responses to experimental conditions across a set of measures (ERP, ERSP, ITC...)
- Creates *non-overlapping IC partitions* making each IC belong to only one cluster.

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### **EEGLAB Study Clustering strategy**

- 1. Cluster on **multiple measures** (**dipole locations**, scalp maps, spectra, ERPs, ITCs, ERSPs, ...) **in one or more conditions**.
- 2. Reduce the dimension of each measure to a principal component subspace.
- 3. Compose a PCA-reduced **position vector** for each component.
- 4. Cluster the composed component vectors using k-means or other.
- Use the computed component measures (not PCA-reduced) to visualize the activities and spatial properties of the clustered components.
- 6. Compute and visualize the **cluster-mean measures**.
- 7. Use clustered Study set data as input into 'std\_???' functions.

## Study IC Clustering





Onton & Makeig, 2007

## EEGLAB Study Clustering procedure



- 1. Identify a set of datasets as an EEGLAB **Study**.
- 2. Specify the **subject** code, subject **group**, **condition** and/or **session** for each dataset in the Study.
- 3. Identify **components to cluster** in each Study dataset.
- 4. Decide on **component measures** to use in clustering the Study and/or to evaluate the obtained component clusters.
- 5. Compute the component measures for each Study dataset.
- 6. Cluster the components on these component measures.
- 7. Review the obtained clusters (e.g., their scalp maps, dipoles, and activity measures).
- 8. Edit the clusters (manually remove/shift components, make subclusters, merge clusters, re-cluster).
- 9. Statistically test differences within or between selected clusters.







statistics within subject and binomial probability between subjects (p < 0.01)

between the two clusters by bootstrap statistics (p < 0.001)



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## **STUDY IC Clustering: Practical Problems**

Large parameter space problem: many different clustering solutions can be produced by changing parameters and measure subsets. Which one should we choose?



N. Bigdely-Shamlo, 2010

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#### Problems with multi-measure clustering In a uniform density distribution,

where are the clusters by location?



## **Problems with multi-measure clustering**

What are the clusters according to location?

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## **Problems with multi-measure clustering**

What are the clusters according to size ?



## **Problems with multi-measure clustering**

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What are the clusters according to location and size?

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Well, it depends on how much weight we give each





- With either clustering method, we basically mix together distances for a subset of EEG measures (ERP, ERSP, ITC, mean spectrum, dipole location).
- This may make clustering distance less interpretable.



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## **Study IC Clustering by Measure Projection**

- Instead, we can directly work on pair-wise similarity matrices and prevent ICs with similarities less than certain threshold (e.g., ERSP corr. < 0.5) to be clustered together.</li>
- The most important measure is **equivalent dipole location**.
- Assuming a certain variability estimate for dipole location (due to error in localization and subject variability), one can also estimate an optimum number of clusters.

#### **Measure Projection asks:**

- 1. Where in 'template brain space' does our data have evidence that our measure of interest is consistent across nearby ICs?
- 2. Which such brain space voxel *domains* show consistent differences?

#### Project Target ERSPs on Equivalent Dipole Locations Measure Projection: RSVP Task Example





N. Bigdely-Shamlo, 2011

**ERSP** Dissimilarity





# **Questions?**

