

Topological Data Analysis for Brain Networks

Relating Functional Brain Network Topology to Clinical Measures of Behavior

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Joint work with

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Big Picture

Goal: Quantify the relationship between brain functional networks and behavioral measures.

Our Contribution: Use topological features based on persistent homology.

Result: Combining correlations with topological features gives better prediction of autism severity than using correlations alone.



functional network



behavior

About Autism Spectrum Disorders (ASD):

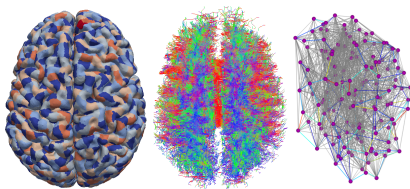
- No cure, causes unknown
- Diagnosis:
 - No systematic method
 - ADOS (Autism Diagnostic Observation Schedule)

Correlate functional brain network to ADOS scores

- Early diagnosis
- Treatment tracking

What is a Brain Network?

- Represents brain regions and pairwise associations
- Computation of Correlation Matrices:
 - Resting state functional MRI (R-fMRI)
 - Preprocessing
 - Define regions of interest (ROIs)
 - Estimate time series signals
 - Compute pairwise associations - Pearson Correlation

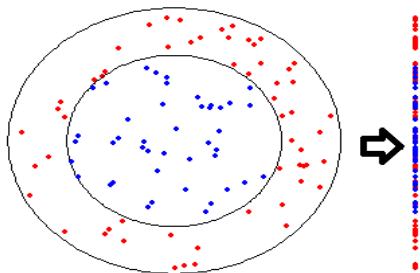


Why Topology?

How to use this data?

- Graph and graph theoretic measures (e.g. small worldness)
 - Require binary associations (thresholding)
- Correlations as features
 - High dimensionality, not enough samples
- Dimensionality reduction: PCA, random projections
 - May lose structures in higher dimensions

Projection - may lose structures in higher dimensions

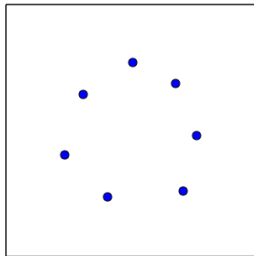
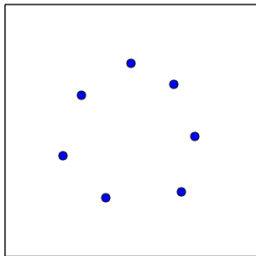


Topology captures structure

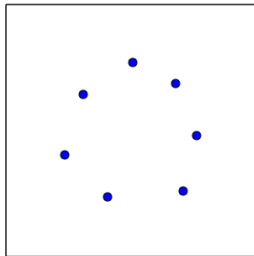
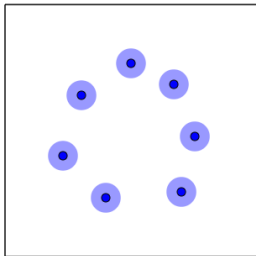
- In higher dimensions
- Across all continuous thresholds

- What are topological features? Homological features:
 - Dim 0 - Connected Components
 - Dim 1 - Tunnels / Loops
 - Dim 2 - Voids
- How to compute them (in a nutshell)?
 - Begin with point cloud
 - Grow balls of diameter t around each point
 - Track features of the union of balls as t increases

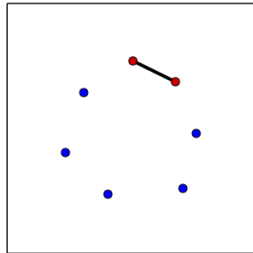
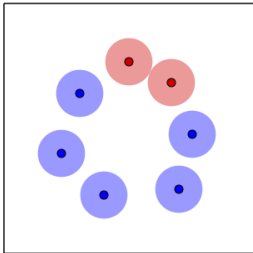
Persistent Homology



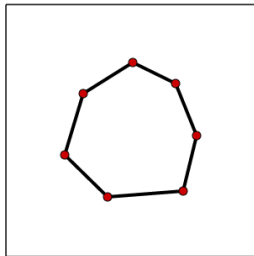
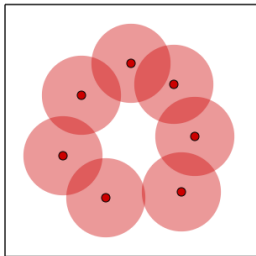
Persistent Homology



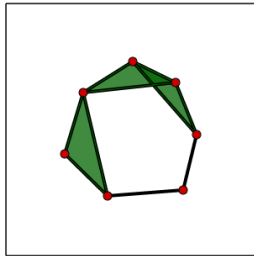
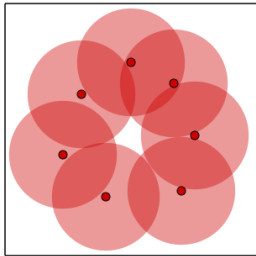
Persistent Homology



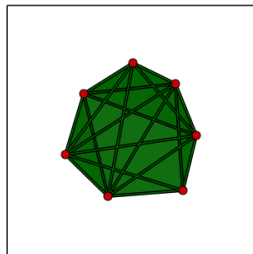
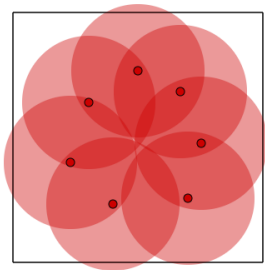
Persistent Homology



Persistent Homology



Persistent Homology



Persistence Diagrams

Persistent homological features - encoded as barcodes or persistent diagrams

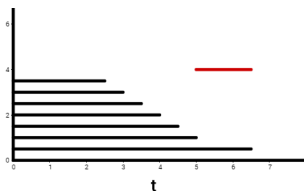


Figure: Barcode

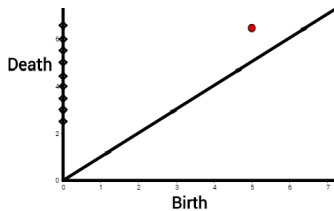
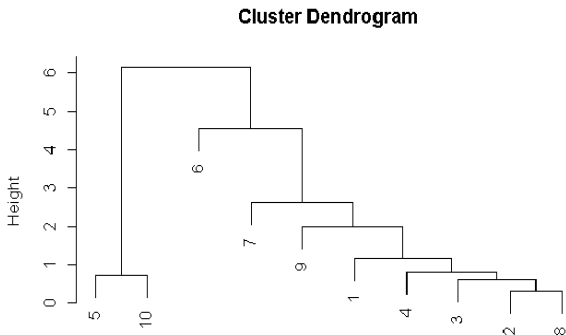


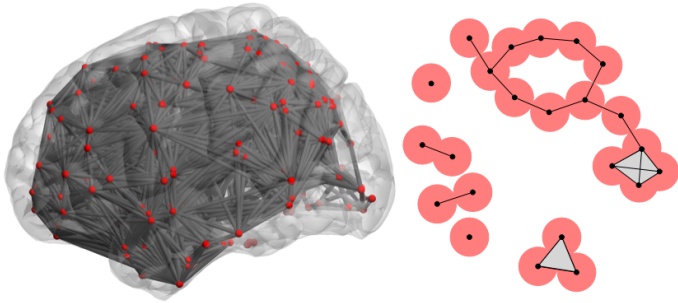
Figure: Persistence Diagram

Interpretation of Connected Components

- Dim 0 features - hierarchical clustering



Computing Topological Features for Brain Networks



Partial Least Squares (PLS) Regression

A dimensionality reduction technique that finds two sets of latent dimensions from datasets X and Y such that their **projections** on the latent dimensions are **maximally co-varying**.

- X - features from brain imaging: correlations, topological features (zero mean)
- Y - clinical measure of behavior: ADOS scores (zero mean)

PLS models the relations between X and Y by means of **score vectors**.

PLS Regression

- n - number of data points
- X - predictor/regressor ($n \times N$), Y - response ($n \times M$)
- PLS - decompose X , Y such that:

$$X = TP^T + E$$

$$Y = UQ^T + F$$

Where

- T , U - latent variables/score vectors ($n \times p$), factor matrices
- P ($N \times p$), Q ($M \times p$) - orthogonal loading matrices
- E ($n \times N$), F ($n \times M$) - residuals/errors
- T , U are chosen such that projections of X , Y , that is, T and U , are maximally co-varying.

PLS Regression: the Algorithm

Iterative NIPALS¹ algorithm

- Find first latent dimension
i.e. find vectors w, c such that

$$t = Xw, \quad u = Yc$$

have maximal covariance

- Deflate previous latent dimensions from X, Y and repeat

¹Nonlinear iterative partial least squares; [Wold 1975]

Kernel form of NIPALS algorithm (kPLS)

1. Initialize random vector u
2. Repeat until convergence
 - (a) $t = Ku/\|Ku\|$
 - (b) $c = Y^T t$
 - (c) $u = Yc/\|Yc\|$
3. Deflate $K = (I - tt^T)K(I - tt^T)$
4. Repeat to compute subsequent latent dimensions

- 87 Subjects: 30 Control, 57 ASD
- ADOS scores: 0 to 21
- 264 ROIs (Power regions)
- 264×264 correlation matrix.
- 34,716 distinct pairwise correlations per subject.

Experiments

- Given: Correlation matrices
- Map to metric space

$$d(x, y) = \sqrt{1 - \text{Cor}(x, y)}$$

- Compute persistence diagrams
- Define inner product of persistence diagrams² (i.e. kernel):
Given two persistence diagrams F, G

$$k_{\sigma}(F, G) = \frac{1}{8\pi\sigma} \sum_{p \in F} \sum_{q \in G} e^{-\frac{\|p-q\|^2}{8\sigma}} - e^{-\frac{\|p-\bar{q}\|^2}{8\sigma}}$$

where for every $q = (x, y) \in G$, $\bar{q} = (y, x)$

²[Reininghaus Huber Bauer Kwitt 2015].

Performed experiments with 3 kernels:

1. K^{Cor} - Euclidean dot product of vectorized correlations
2. $K^{TDA} = w_0 K^{TDA_0} + (1 - w_0) K^{TDA_1}$
 - K^{TDA_0} - using only Dim 0 features
 - K^{TDA_1} - using only Dim 1 features
3. $K^{TDA+Cor} = w_0 K^{TDA_0} + w_1 K^{TDA_1} + (1 - w_0 - w_1) K^{Cor}$

Baseline predictor - mean ADOS score

Experiments

- Leave one out cross validation over parameters
 - σ_0, σ_1 - ($\log_{10} \sigma$) from -8.0 to 6.0 by 0.2
 - w_0, w_1 - from 0.0 to 1.0 by 0.05
- k^{TDA} parameters: $\sigma_0 = -6.6, \sigma_1 = 1.8, w_1 = 0.95$
- $k^{TDA+Cor}$ parameters: $\sigma_0 = -7.8, \sigma_1 = 2.8, w_0 = 0.1, w_1 = 0.4$
- Compute RMSE
- Permutation test for significance

Results

	RMSE	ADOS mean	K^{TDA}	K^{cor}
ADOS mean	6.4302	-	-	-
K^{TDA}	6.3553	0.316	-	-
K^{cor}	6.0371	0.055	0.095	-
$K^{TDA+cor}$	6.0156	0.048	0.075	0.288

Table: ADOS prediction results. Columns 2 to 4 are p-values for the permutation test of improvement of row method over column method.

Result Highlights:

- Baseline RMSE: 6.4302
- $K^{TDA+Cor}$:
 - Only method statistically significant over baseline
 - Permutation test p-value: 0.048
 - RMSE: 6.0156

Conclusion

- Augmenting correlations with topological features gives a **better** prediction of autism severity than using correlations alone
- (Hopefully) topological features derived from R-fMRI have the **potential** to explain the connection between functional brain networks and autism severity

Many things to try

- Alternatives to correlation
- Different distance metric
- Different kernel
- Multi-site data
- Classification (combine with TDA features)

Kernel Partial Least Squares Regression for Relating Functional Brain Network Topology to Clinical Measures of Behavior

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Thank you!

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functional network



behavior

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