# Learning Joint and Individual Structure in Network Data with Covariates

Difussion MRI brain connectomes from the



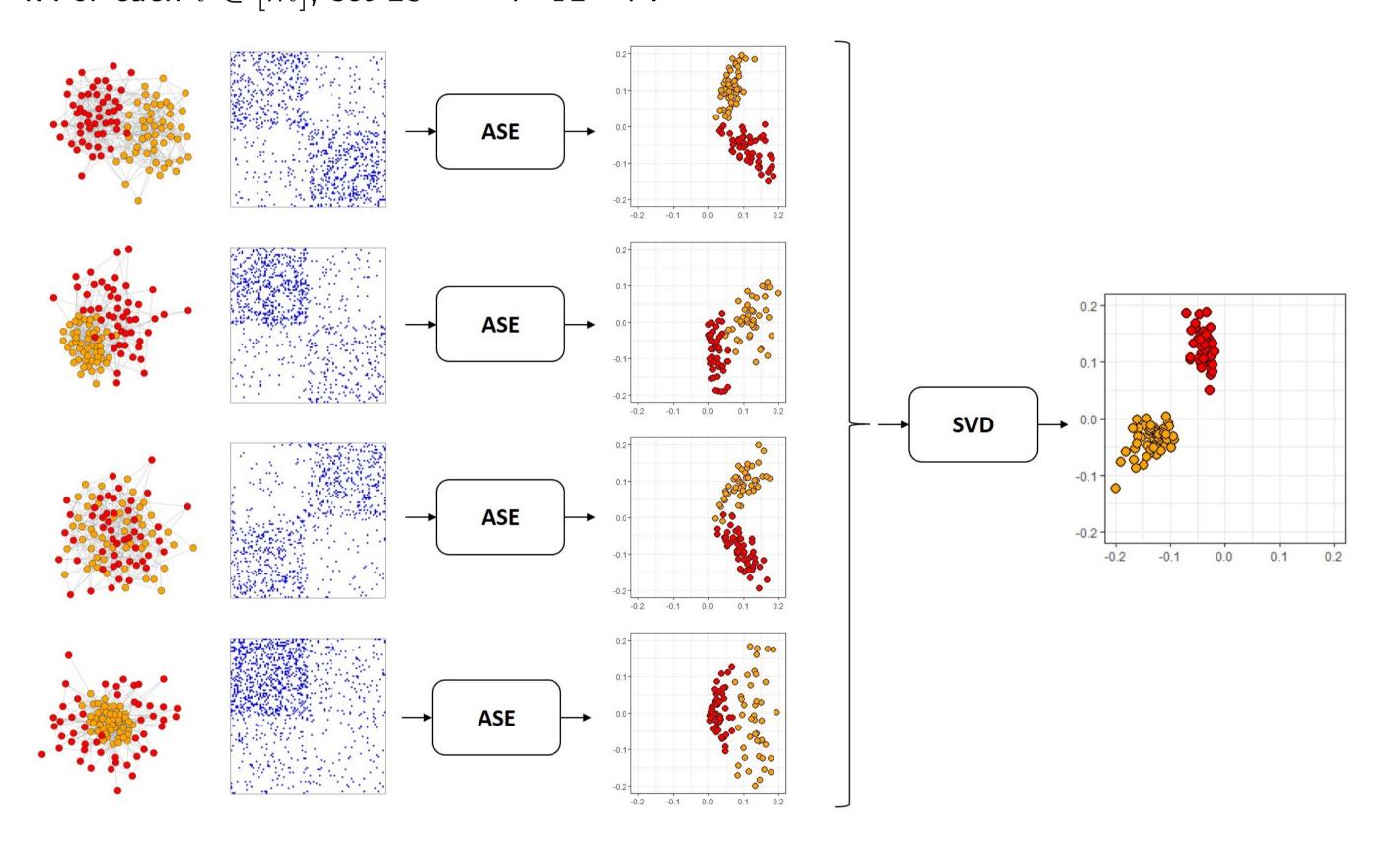
## Modeling multiple heterogeneous networks

#### **Motivation:**

- We present the *common subspace independent-edge* (COSIE) model which describes a collection of networks with a shared latent structure on the vertices but potentially different connectivity patterns for each graph [1].
- COSIE encompasses many other popular network representations, including the stochastic blockmodel
- A joint spectral embedding the *multiple adjacency spectral embedding*-leads to consistent estimation that is computationally efficient
- MASE estimates yield state-of-the-art performance on subsequent inference tasks, including dimensionality reduction, hypothesis testing and community detection
- Models for multiple network data are critical in statistical network theory and across multiple domains, including neuroscience, biology and the social sciences.
- Challenges in modeling graph differences while retaining sufficient model simplicity to render estimation feasible.
- Existing models require strong assumptions that limit their flexibility or scalability

# Multiple adjacency spectral embedding (MASE)

- 1. For each  $i \in [m]$ , obtain the d-dimensional unscaled adjacency spectral embedding of  $\mathbf{A}^{(i)}$ , denoted by  $\hat{\mathbf{V}} \in \mathbb{R}^{n \times d}$  and corresponding to the d leading eigenvectors
- 2. Let  $\hat{\mathbf{U}} = (\hat{\mathbf{V}}^{(1)} \cdots \hat{\mathbf{V}}^{(m)})$  be the  $n \times (md)$  matrix of concatenated ASEs.
- 3. Let  $\hat{\mathbf{V}} \in \mathbb{R}^{n \times d}$  be the matrix containing the d leading left singular values of  $\hat{\mathbf{U}}$ .
- 4. For each  $i \in [m]$ , set  $\hat{\mathbf{R}}^{(i)} = \hat{\mathbf{V}}^T \mathbf{A}^{(i)} \hat{\mathbf{V}}$ .



# Common subspace independent edge (COSIE) model

- ullet Consider a sample of m graphs with n labeled nodes.
- ullet Denote the graphs by their adjacency matrices  $\mathbf{A}^{(1)},\ldots,\mathbf{A}^{(m)}\in\{0,1\}^{n imes n}$
- ullet Each graph  $\mathbf{A}^{(i)}$  is modeled as independent-edge with parameter  $\mathbf{P}^{(i)} \in \mathbb{R}^{n imes n}$

$$\mathbf{A}_{uv}^{(i)} \sim \mathrm{Ber}(P_{uv}^{(i))}).$$

#### **COSIE** model

• The sample of graphs is jointly distributed according to the COSIE model if

$$\mathbf{P}^{(i)} := \mathbf{V} \mathbf{R}^{(i)} \mathbf{V}^T$$
.

- ullet  $\mathbf{V} \in \mathbb{R}^{n \times d}$  is a matrix with orthogonal columns, with its rows representing vertex latent positions
- $ullet \mathbf{R}^{(i)} \in \mathbb{R}^{d \times d}$  is a score matrix, potentially different for each graph
- ullet The parameter d controls the complexity of the model

# Theoretical properties

### Consistency of common invariant subspace estimator

 $\bullet$  Under some assumptions on the smallest eigenvalue of the score matrices and on the sparsity of the graphs, the estimate of  ${\bf V}$  obtained by MASE satisfies

$$\mathbb{E}\left[\min_{\mathbf{W}\in\mathcal{O}_d}\left\|\hat{\mathbf{V}}-\mathbf{V}\mathbf{W}\right\|\right]\lesssim \sqrt{\frac{d}{mn}}+\frac{\sqrt{d}}{n}.$$

• When V has only d different rows, COSIE is equivalent to the stochastic blockmodel, and k-means clustering expected error in community detection is  $O\left(\sqrt{\frac{d}{m}} + \frac{1}{\sqrt{n}}\right)$ .

## Asymptotic normality of the score matrices

ullet The entries of the estimated score matrices  $\hat{\mathbf{R}}^{(i)}$  are asymptotically normally distributed, in particular, as the size of the graphs n increases

$$\frac{1}{\sigma_{ijk}} \left( \hat{\mathbf{R}}^{(i)} - \mathbf{W} \mathbf{R}^{(i)} \mathbf{W}^T + \mathbf{H} \right)_{jk} \xrightarrow{d} \mathcal{N}(0, 1)$$

where  $\sigma_{ijk} = O(1)$ ,  $\mathbb{E}[\|\mathbf{H}\|_F] = O(d/\sqrt{m})$  and  $\|\mathbf{R}^{(i)}\|_F \to \infty$ .

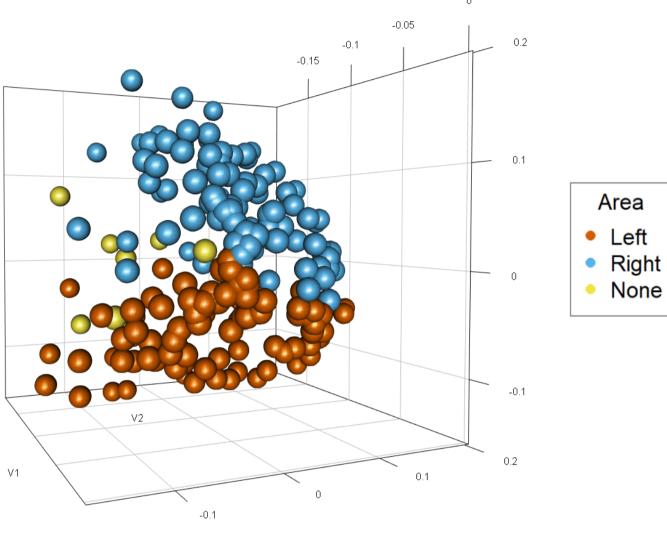
## Brain network analysis

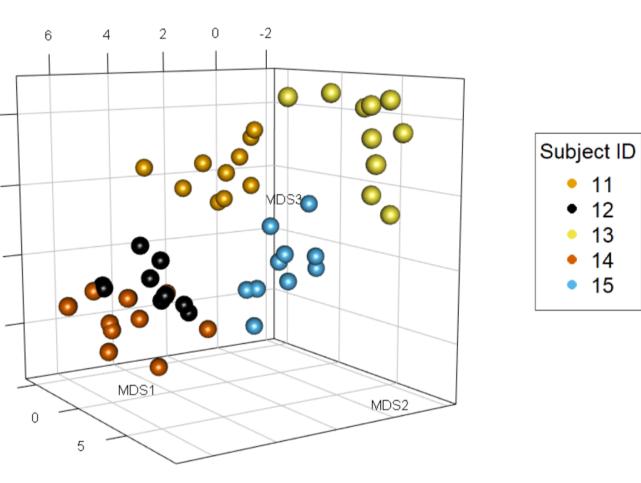
#### HNU1 data

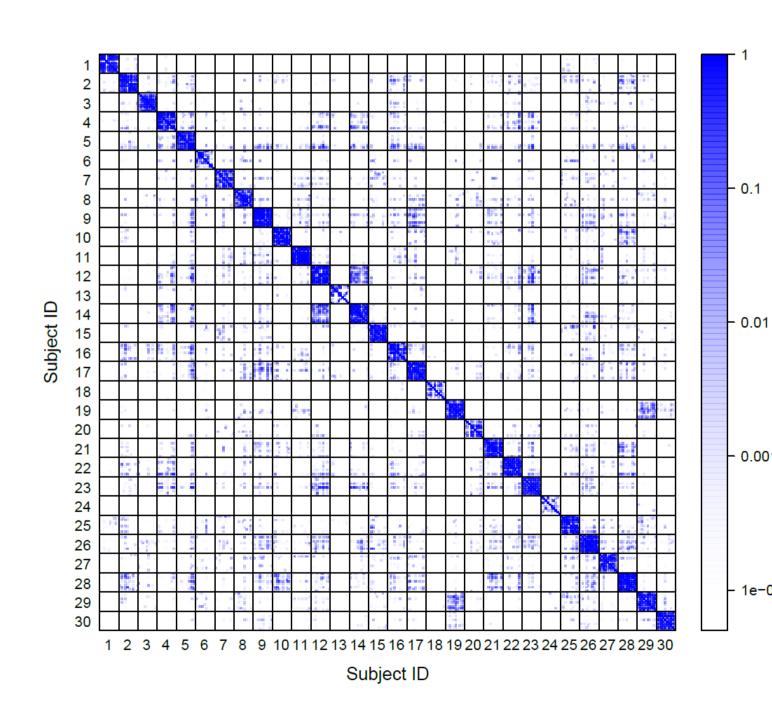
- 300 graphs constructed from diffusion magnetic resonance imaging (dMRI), with n=200 nodes
- 30 different healthy subjects, and 10 graphs per subject
- **Goal:** identify differences and similarities between graphs

#### Dimensionality reduction

- ullet The vertex embedding  $\hat{\mathbf{V}}$  obtained by MASE (top figure) reflects the anatomical location of the vertices in the brain.
- A multidimensional scaling of the distance between the score matrices  $\{\mathbf{R}^{(i)}\}_{i=1}^{300}$  (bottom figure) puts graphs from the same subject closer to each other







Matrix of p-values for the equal distribution test of every pair of graphs

## Graph hypothesis testing

- For each pair of graphs i and j, we test the hypothesis that their distribution is the same, i.e.,  $H_0: \mathbf{R}^{(i)} = \mathbf{R}^{(j)}$
- ullet Test statistic  $\|\hat{\mathbf{R}}^{(i)} \hat{\mathbf{R}}^{(j)}\|_F$
- Semiparametric bootstrap: estimate the parameters with MASE to sample new graphs
- The p-values of the test (left figure) are generally high for pairs of graphs from the same subject (diagonal entries) and low for different subjects

#### Acknowledgements

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#### References

[1] Jesús Arroyo, Avanti Athreya, Joshua Cape, Guodong Chen, Carey E Priebe, and Joshua T Vogelstein, "Inference for multiple heterogeneous networks with a common invariant subspace," arXiv:1906.10026, 2019.

Open R source code for MASE is available at https://github.com/jesusdaniel/mase, and in the Python GraSPy package at https://neurodata.io/graspy.