

Classification of gene expression data with gene networks

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- 1 Motivation
- 2 Using gene networks as prior knowledge
- 3 Application
- 4 Conclusion

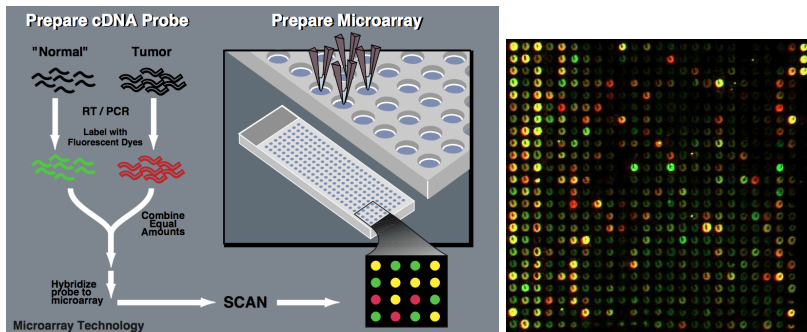
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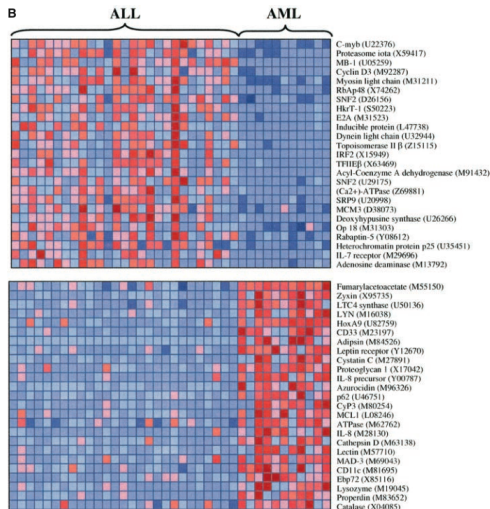
Tissue profiling with DNA chips



Data

- Gene expression measures for **more than 10k genes**
- Measured typically on **less than 100 samples** of two (or more) different classes (e.g., different tumors)

Tissue classification from microarray data



Goal

- Design a **classifier** to automatically assign a class to future samples from their expression profile
- **Interpret** biologically the differences between the classes

The approach

- Each sample is represented by a vector $x = (x_1, \dots, x_p)$ where $p > 10^5$ is the number of probes
- **Classification**: given the set of labeled sample, learn a linear decision function:

$$f_{\beta}(x) = \sum_{i=1}^p \beta_i x_i + \beta_0 ,$$

that is positive for one class, negative for the other

- **Interpretation**: the weight β_i quantifies the influence of gene i for the classification

Empirical risk minimization

Estimate the weights β_i by **minimizing an empirical error** on the training set:

$$\min_{\beta \in \mathbb{R}^{p+1}} \frac{1}{n} \sum_{i=1}^n l(f_{\beta}(x_i), y_i),$$

where $l(y, f(x))$ is a loss function.

Pitfalls

- **Statistics does not apply** (?): 100 samples in 10^5 dimensions!
- It is necessary to **reduce the complexity** of the problem with **prior knowledge**.

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Example : Norm Constraints

The approach

A common method in statistics to learn with few samples in high dimension is to **constrain the Euclidean norm of β**

$$\|\beta\|_2^2 = \sum_{i=1}^p \beta_i^2,$$

(ridge regression, support vector machines...)

Pros

- Good performance in classification

Cons

- Limited interpretation (small weights)
- No prior biological knowledge

Example : Feature Selection

The approach

Constrain most weights to be 0, i.e., **select a few genes** (< 100) whose expression are enough for classification. Interpretation is then about the selected genes. Examples:

- Greedy feature selection (T-tests, ...)
- Constrain the norm of β : **LASSO** penalty ($\|\beta\|_1 = \sum_{i=1}^p |\beta_i|$), **elastic net** penalty ($\|\beta\|_1 + \|\beta\|_2$), ...)

Pros

- Good performance in classification
- **Biomarker** selection
- Interpretability

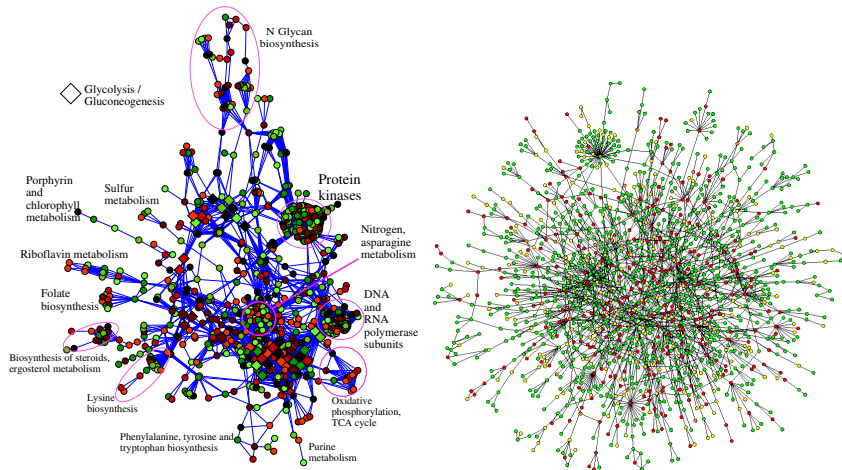
Cons

- The gene selection process is usually **not robust**
- No use of prior biological knowledge

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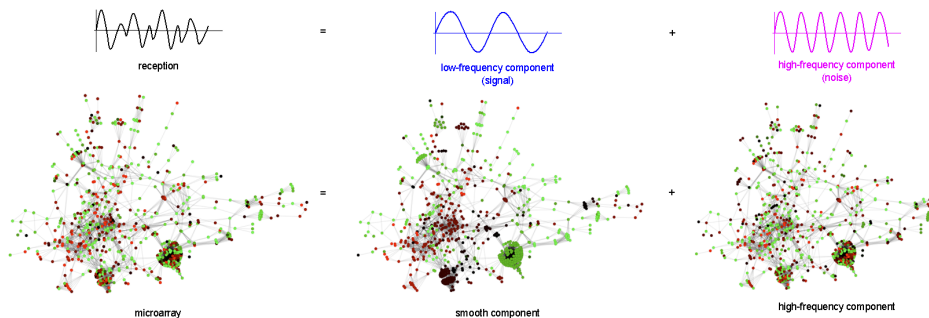
Gene networks



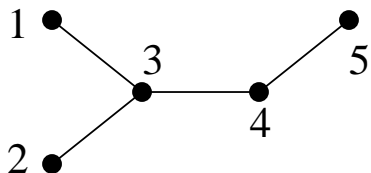
Motivation

- Basic biological functions usually involve the **coordinated action of several proteins**:
 - Formation of **protein complexes**
 - Activation of metabolic, signalling or regulatory **pathways**
- Many pathways and protein-protein interactions are **already known**
- **Hypothesis**: the weights of the classifier should be “coherent” with respect to this **prior knowledge**

The idea



- 1 Use the gene network to extract the “important information” in gene expression profiles by **Fourier analysis** on the graph
- 2 Learn a linear classifier on the **smooth components**

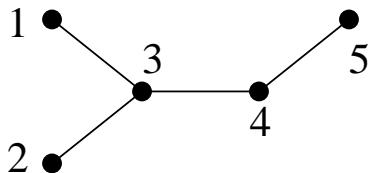


$$A = \begin{pmatrix} 0 & 0 & 1 & 0 & 0 \\ 0 & 0 & 1 & 0 & 0 \\ 1 & 1 & 0 & 1 & 0 \\ 0 & 0 & 1 & 0 & 1 \\ 0 & 0 & 0 & 1 & 0 \end{pmatrix}, \quad D = \begin{pmatrix} 1 & 0 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 & 0 \\ 0 & 0 & 3 & 0 & 0 \\ 0 & 0 & 0 & 2 & 0 \\ 0 & 0 & 0 & 0 & 1 \end{pmatrix}$$

Graph Laplacian

Definition

The Laplacian of the graph is the matrix $L = D - A$.



$$L = D - A = \begin{pmatrix} 1 & 0 & -1 & 0 & 0 \\ 0 & 1 & -1 & 0 & 0 \\ -1 & -1 & 3 & -1 & 0 \\ 0 & 0 & -1 & 2 & -1 \\ 0 & 0 & 0 & 1 & 1 \end{pmatrix}$$

Lemma

Let $L = D - A$ be the Laplacian of the graph:

- For any $f : \mathcal{X} \rightarrow \mathbb{R}$,

$$f^T L f = \sum_{i \sim j} (f(\mathbf{x}_i) - f(\mathbf{x}_j))^2$$

- L is a **symmetric positive semi-definite** matrix
- 0 is an **eigenvalue** with multiplicity equal to the number of connected components.

Proof: link between $\Omega(f)$ and L

$$\begin{aligned}\sum_{i \sim j} (f(\mathbf{x}_i) - f(\mathbf{x}_j))^2 &= \sum_{i \sim j} \left(f(\mathbf{x}_i)^2 + f(\mathbf{x}_j)^2 - 2f(\mathbf{x}_i)f(\mathbf{x}_j) \right) \\ &= \sum_{i=1}^m D_{i,i} f(\mathbf{x}_i)^2 - 2 \sum_{i \sim j} f(\mathbf{x}_i) f(\mathbf{x}_j) \\ &= \mathbf{f}^\top D \mathbf{f} - \mathbf{f}^\top A \mathbf{f} \\ &= \mathbf{f}^\top L \mathbf{f}\end{aligned}$$

Proof: eigenstructure of L

- L is symmetric because A and D are symmetric.
- For any $f \in \mathbb{R}^m$, $f^\top Lf \geq 0$, therefore the (real-valued) eigenvalues of L are ≥ 0 : L is therefore positive semi-definite.
- f is an eigenvector associated to eigenvalue 0
 - iff $f^\top Lf = 0$
 - iff $\sum_{i \sim j} (f(\mathbf{x}_i) - f(\mathbf{x}_j))^2 = 0$,
 - iff $f(\mathbf{x}_i) = f(\mathbf{x}_j)$ when $i \sim j$,
 - iff f is constant (because the graph is connected).

Definition

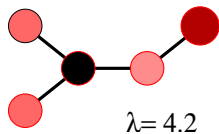
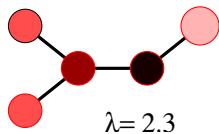
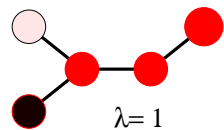
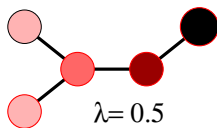
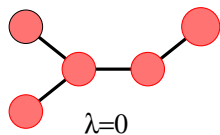
- The **eigenvectors** e_1, \dots, e_n of L with eigenvalues $0 = \lambda_1 \leq \dots \leq \lambda_n$ form a basis called **Fourier basis**
- For any $f : V \rightarrow \mathbb{R}$, the **Fourier transform** of f is the vector $\hat{f} \in \mathbb{R}^n$ defined by:

$$\hat{f}_i = f^\top e_i, \quad i = 1, \dots, n.$$

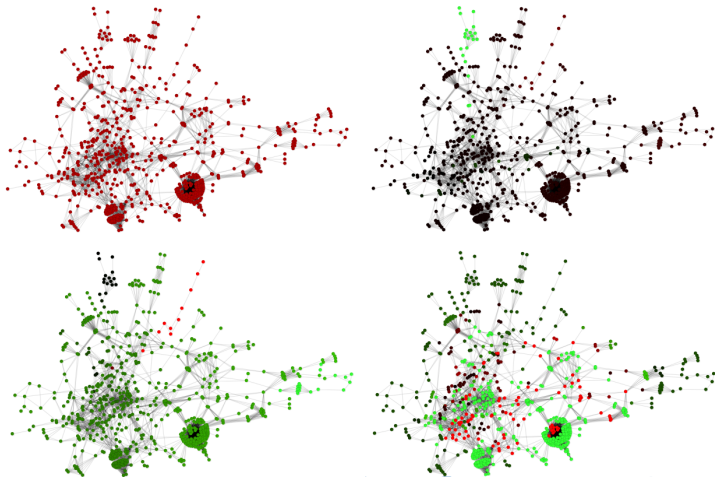
- Obviously the **inverse Fourier formula** holds:

$$f = \sum_{i=1}^n \hat{f}_i e_i.$$

Fourier basis



Fourier basis



Definition

- Let $\phi : \mathbb{R}^+ \rightarrow \mathbb{R}^+$ be **non-increasing**.
- A smoothing operator S_ϕ transform a function $f : V \rightarrow \mathbb{R}$ into a smoothed version:

$$S_\phi(f) = \sum_{i=1}^n \hat{f}_i \phi(\lambda_i) e_i.$$

Examples

- Identity operator ($S_\phi(f) = f$):

$$\phi(\lambda) = 1, \quad \forall \lambda$$

- Low-pass filter:

$$\phi(\lambda) = \begin{cases} 1 & \text{if } \lambda \leq \lambda^*, \\ 0 & \text{otherwise.} \end{cases}$$

- Attenuation of high frequencies:

$$\phi(\lambda) = \exp(-\beta\lambda).$$

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Working with smoothed profiles

- Classical methods for linear classification and regression with a ridge penalty solve:

$$\min_{\beta \in \mathbb{R}^p} \frac{1}{n} \sum_{i=1}^n l(\beta^\top f_i, y_i) + \lambda \beta^\top \beta.$$

- Applying these algorithms on the smooth profiles means solving:

$$\min_{\beta \in \mathbb{R}^p} \frac{1}{n} \sum_{i=1}^n l(\beta^\top \mathcal{S}_\phi(f_i), y_i) + \lambda \beta^\top \beta.$$

Smooth solution

Lemma

This is equivalent to:

$$\min_{v \in \mathbb{R}^p} \frac{1}{n} \sum_{i=1}^n l(v^\top f_i, y_i) + \lambda \sum_{i=1}^p \frac{\hat{v}_i^2}{\phi(\lambda_i)},$$

hence the linear classifier v is **smooth**.

Proof

- Let $v = \sum_{i=1}^n \phi(\lambda_i) e_i e_i^\top \beta$, then

$$\beta^\top \mathcal{S}_\phi(f_i) = \beta^\top \sum_{i=1}^n \hat{f}_i \phi(\lambda_i) e_i = f_i^\top v.$$

- Then $\hat{v}_i = \phi(\lambda_i) \hat{\beta}_i$ and $\beta^\top \beta = \sum_{i=1}^n \frac{\hat{v}_i^2}{\phi(\lambda_i)^2}$.

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Smoothing kernel

Kernel methods (SVM, kernel ridge regression..) only need the **inner product between smooth profiles**:

$$\begin{aligned}K(f, g) &= \mathbf{S}_\phi(f)^\top \mathbf{S}_\phi(g) \\&= \sum_{i=1}^n \hat{f}_i \hat{g}_i \phi(\lambda_i)^2 \\&= f^\top \left(\sum_{i=1}^n \phi(\lambda_i)^2 \mathbf{e}_i \mathbf{e}_i^\top \right) g \\&= f^\top \mathbf{K}_\phi g,\end{aligned}\tag{1}$$

with

$$\mathbf{K}_\phi = \sum_{i=1}^n \phi(\lambda_i)^2 \mathbf{e}_i \mathbf{e}_i^\top.$$

Examples

- For $\phi(\lambda) = \exp(-t\lambda)$, we recover the **diffusion kernel**:

$$K_\phi = \exp_M(-2tL).$$

- For $\phi(\lambda) = 1/\sqrt{1+\lambda}$, we obtain

$$K_\phi = (L + I)^{-1},$$

and the penalization is:

$$\sum_{i=1}^n \frac{\hat{v}_i^2}{\phi(\lambda_i)} = \mathbf{v}^\top (L + I) \mathbf{v} = \|\mathbf{v}\|_2^2 + \sum_{i \sim j} (v_i - v_j)^2.$$

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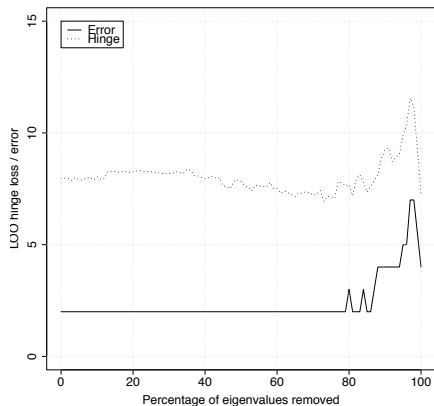
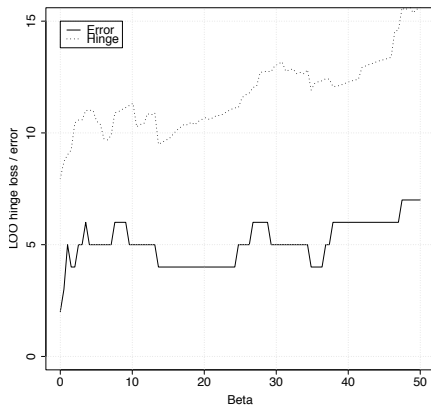
Expression

- Study the effect of low irradiation doses on the yeast
- 12 non irradiated vs 6 irradiated
- Which pathways are involved in the response at the transcriptomic level?

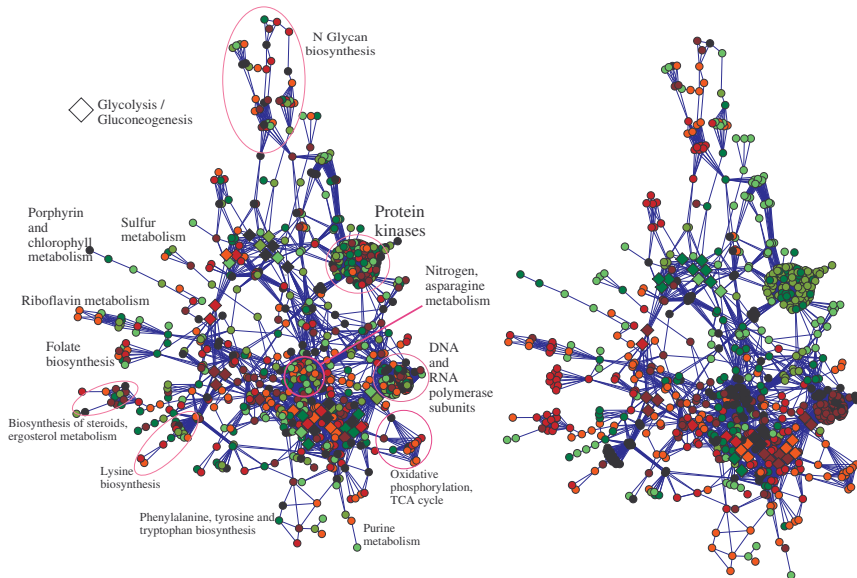
Graph

- KEGG database of metabolic pathways
- Two genes are connected if they code for enzymes that catalyze successive reactions in a pathway (**metabolic gene network**).
- 737 genes, 4694 vertices.

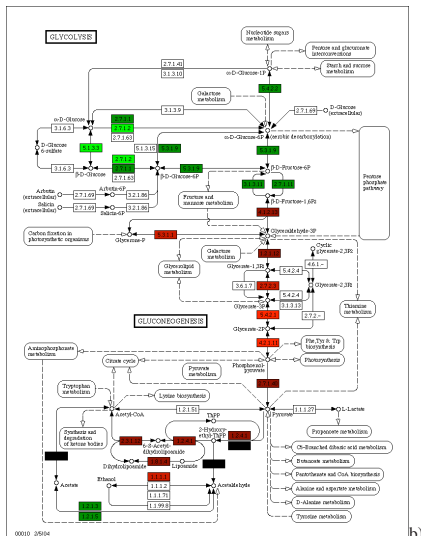
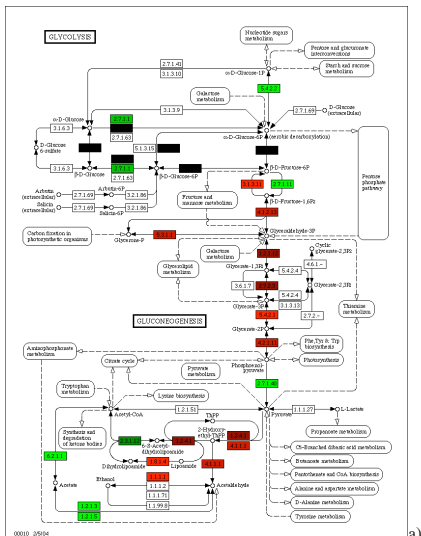
Classification performance



Classifier



Classifier



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Conclusion

- Use the gene graph to encode **prior knowledge** about the classifier.
- Prior knowledge is always needed to classify few examples in large dimensions (sometimes implicitly)
- Future work: validation of the method on more data, other formulations, directed graphs...

Acknowledgements

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- Marie Dutreix (Curie Institute)

Reference

F. Rapaport, A. Zynoviev, M. Dutreix, E. Barillot and J.-P. Vert, Classification of microarray data using gene networks, *BMC Bioinformatics* 8:35, 2007.