

CAP 5510: Introduction to Bioinformatics  
CGS 5166: Bioinformatics Tools

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[www.cis.fiu.edu/~giri/teach/BioinfS11.html](http://www.cis.fiu.edu/~giri/teach/BioinfS11.html)

# In Memoriam



Isabel Melo, Accountant  
Feb 16, 2011

# Gene Expression

- ❑ Process of transcription and/or translation of a gene is called **gene expression**.
- ❑ Every cell of an organism has the same genetic material, but different genes are **expressed** at different times.
- ❑ Patterns of gene expression in a cell is indicative of its state.

# Hybridization

- If two complementary strands of DNA or mRNA are brought together under the right experimental conditions they will hybridize.
- $A$  hybridizes to  $B \Rightarrow$ 
  - $A$  is reverse complementary to  $B$ , or
  - $A$  is reverse complementary to a subsequence of  $B$ .
- It is possible to experimentally verify whether  $A$  hybridizes to  $B$ , by labeling  $A$  or  $B$  with a radioactive or fluorescent tag, followed by excitation by laser.

# Measuring gene expression

- Gene expression for a single gene can be measured by extracting mRNA from the cell and doing a simple **hybridization** experiment.
- Given a sample of cells, gene expression for every gene can be measured using a single microarray experiment.

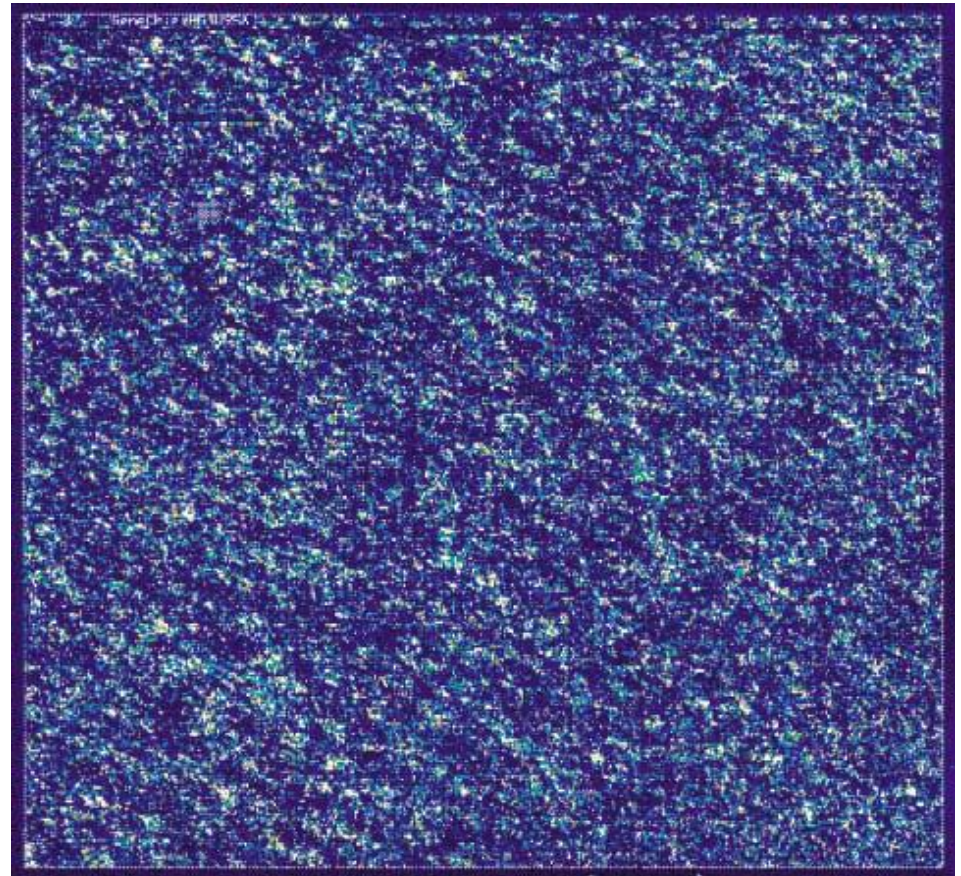
# Microarray/DNA chip technology

- High-throughput method to study gene expression of thousands of genes simultaneously.
- Many applications:
  - Genetic disorders & Mutation/polymorphism detection
  - Study of disease subtypes
  - Drug discovery & toxicology studies
  - Pathogen analysis
  - Differing expressions over time, between tissues, between drugs, across disease states

# Microarray Data

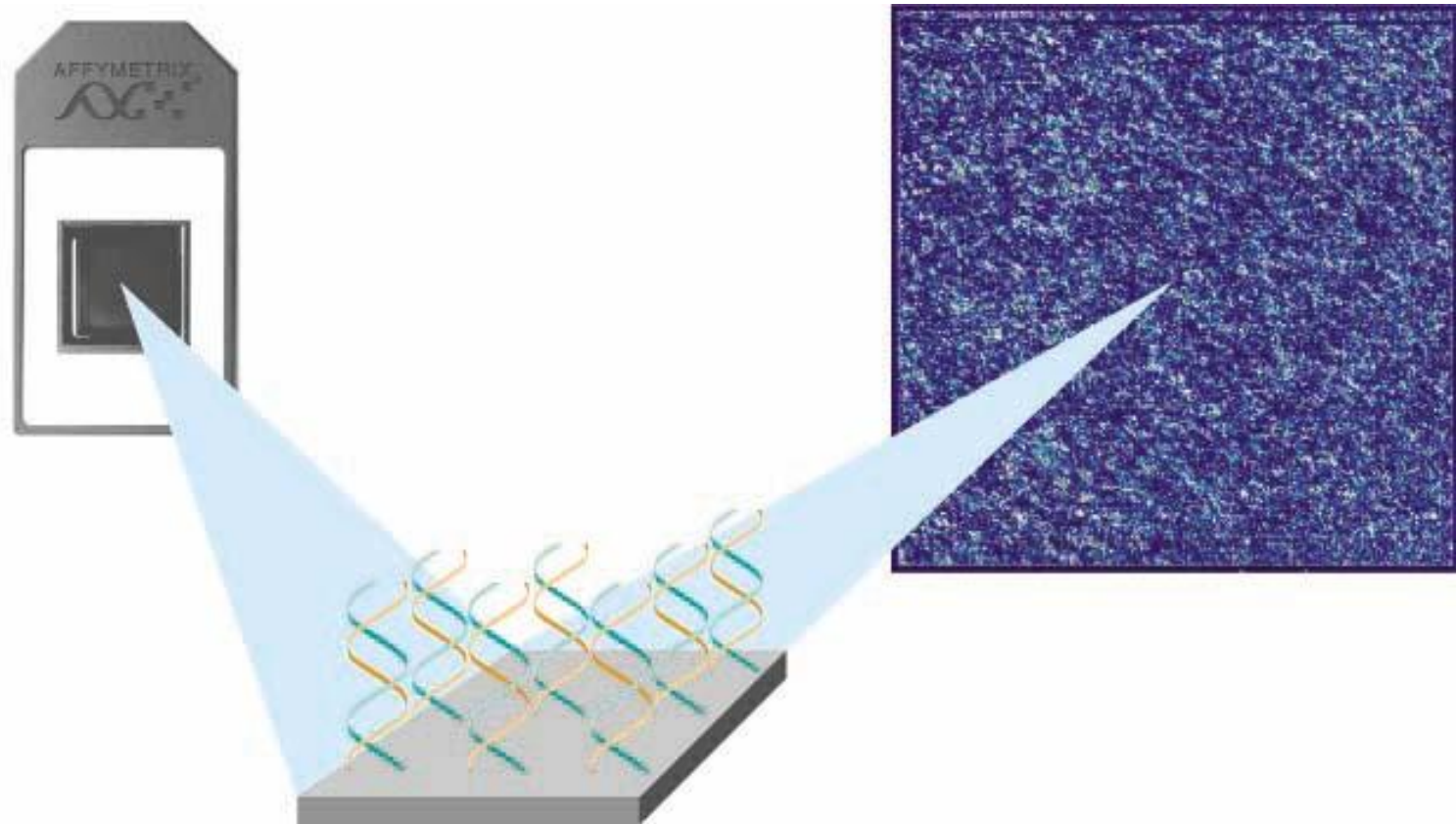
Gene	Expression Level
Gene1	
Gene2	
Gene3	
...	

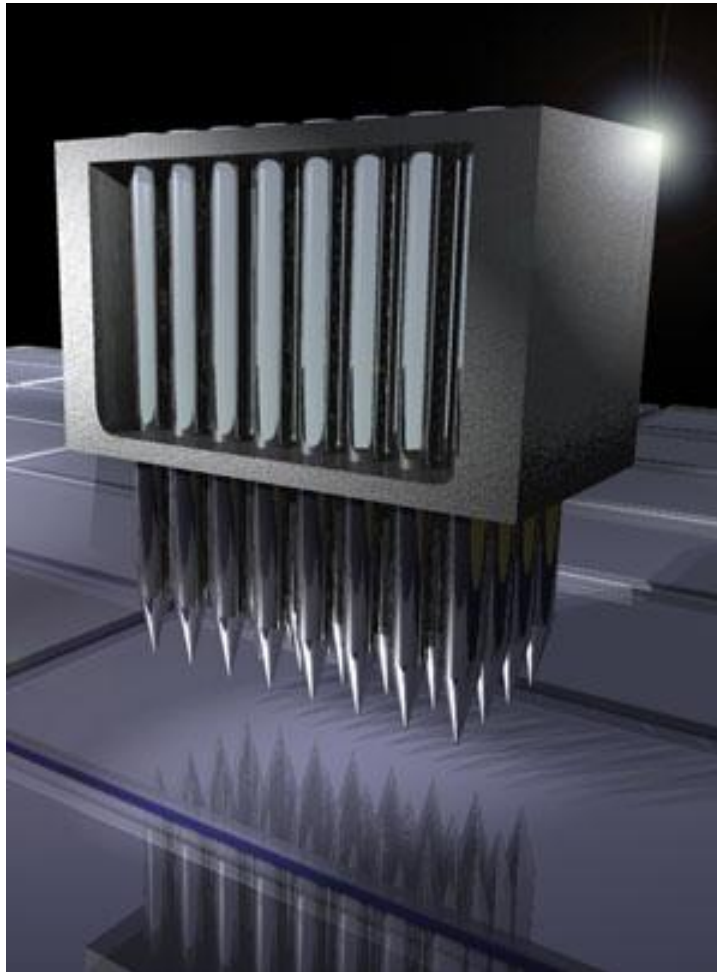
# Gene Chips



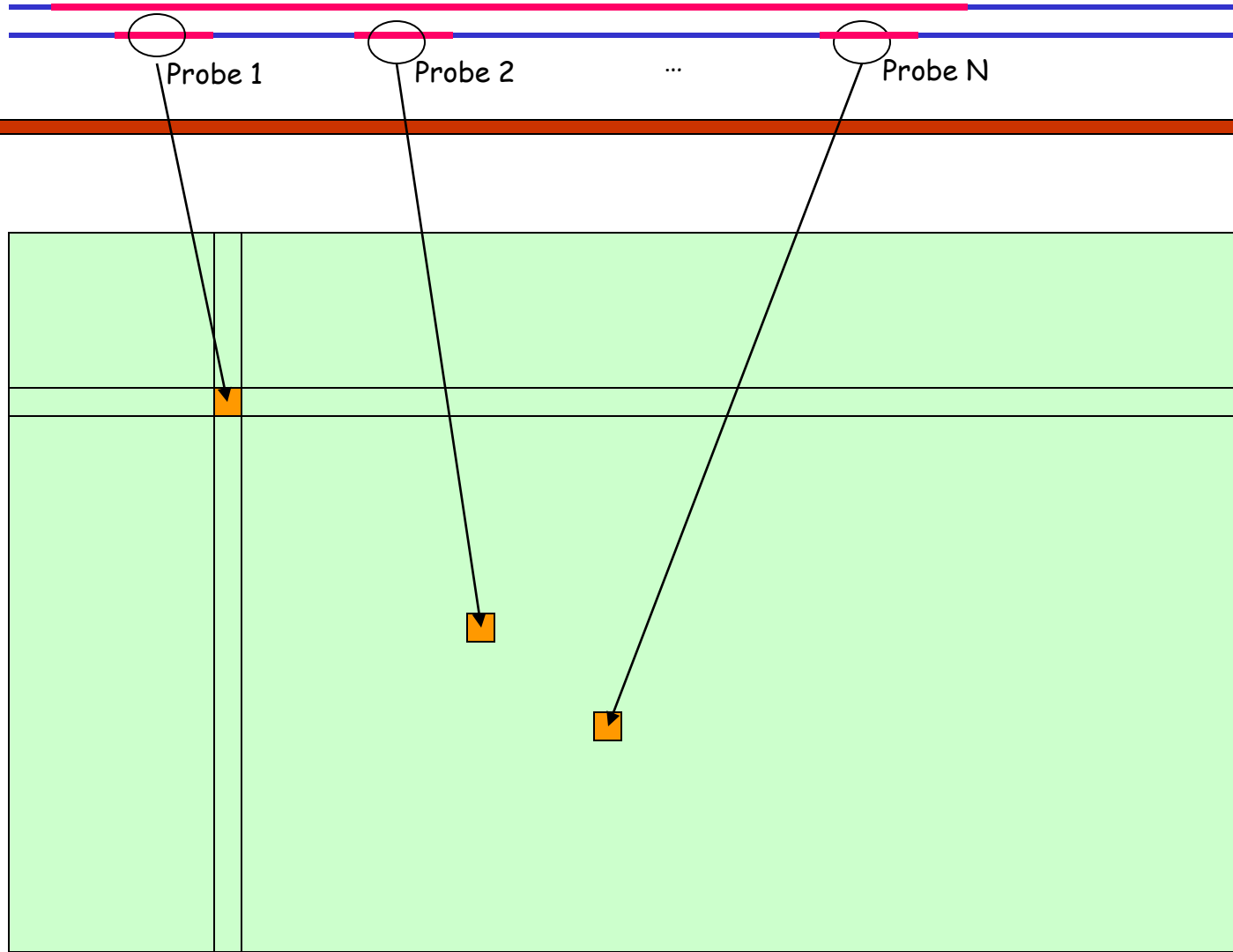


# DNA Chips & Images





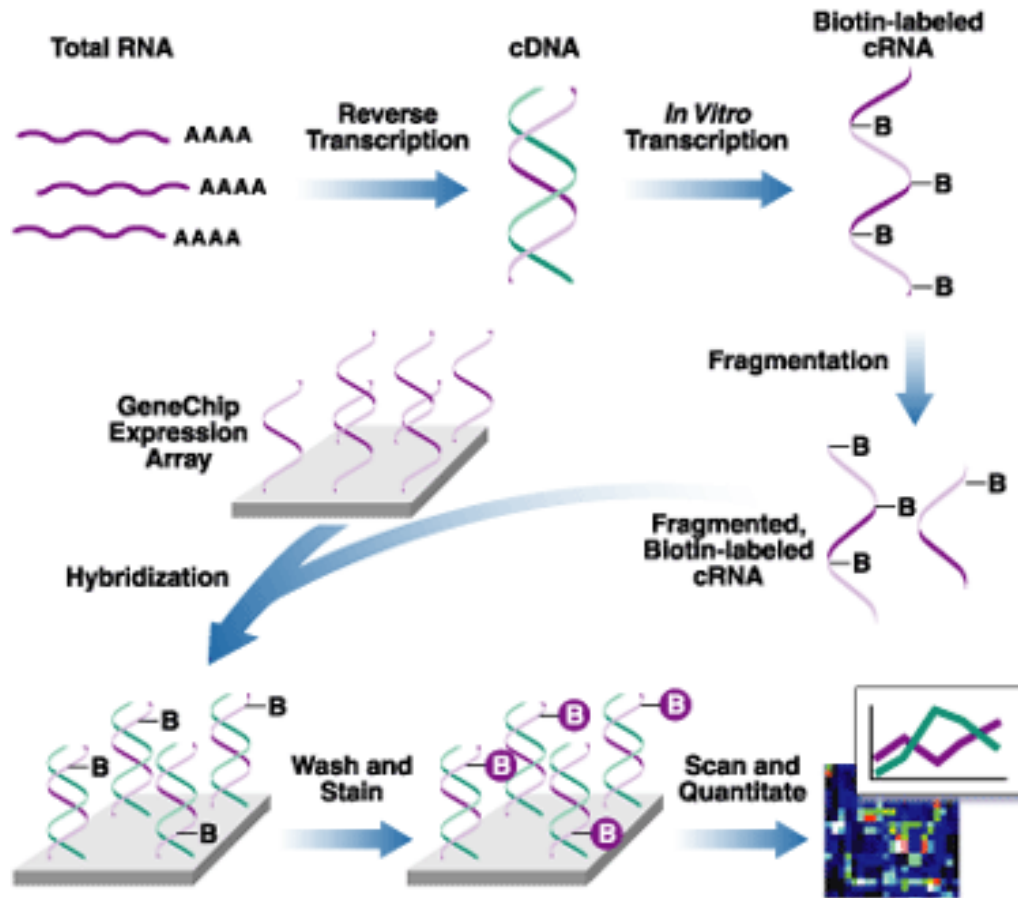
# Gene g



# Microarray/DNA chips (Simplified)

- ❑ Construct **probes** corresponding to reverse complements of genes of interest.
- ❑ Microscopic quantities of probes placed on solid surfaces at defined spots on the chip.
- ❑ Extract mRNA from sample cells and **label** them.
- ❑ Apply labeled sample (mRNA extracted from cells) to every spot, and allow hybridization.
- ❑ Wash off unhybridized material.
- ❑ Use optical detector to measure amount of fluorescence from each spot.

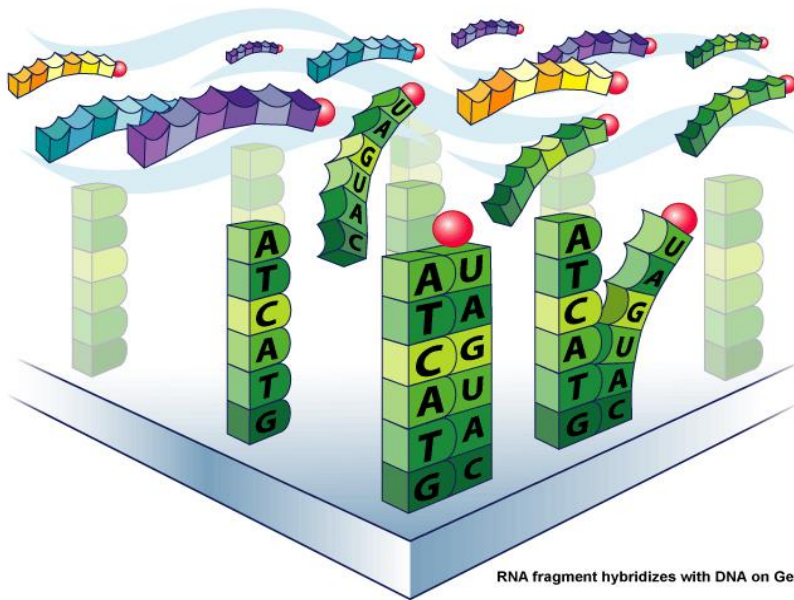
# Affymetrix DNA chip schematic



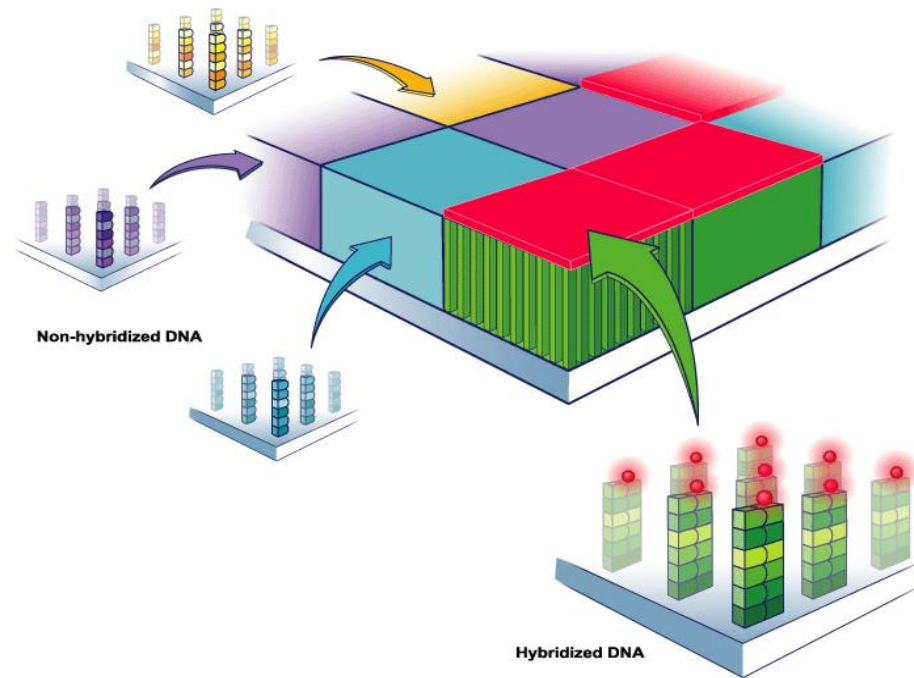
[www.affymetrix.com](http://www.affymetrix.com)

# What's on the slide?

RNA fragments with fluorescent tags from sample to be tested



Shining a laser light at GeneChip® array causes tagged DNA fragments that hybridized to glow



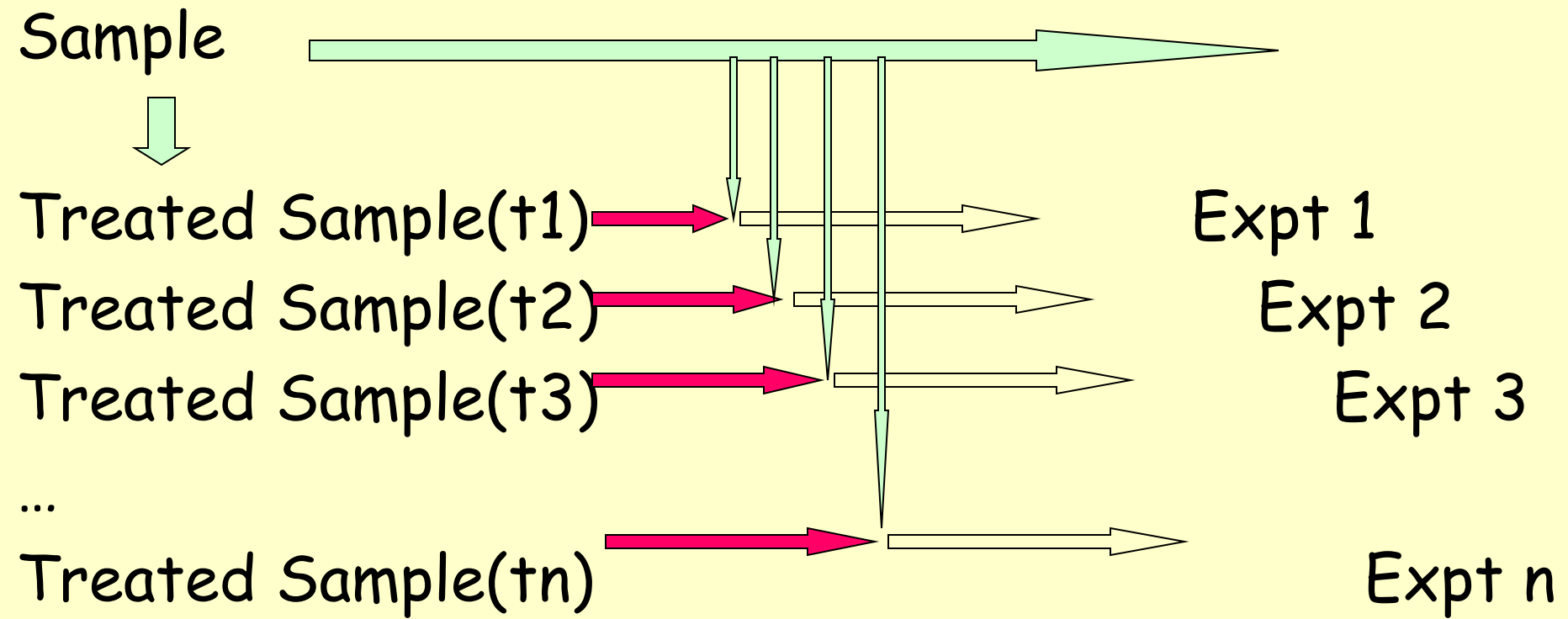
# Microarrays: competing technologies

## □ Affymetrix & Agilent

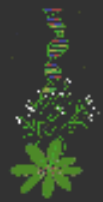
### □ Differ in:

- method to place DNA: Spotting vs. photolithography
- Length of probe
- Complete sequence vs. series of fragments

# Study effect of treatment over time







AFGC

# 2-color DNA microarray



Treated

mRNA

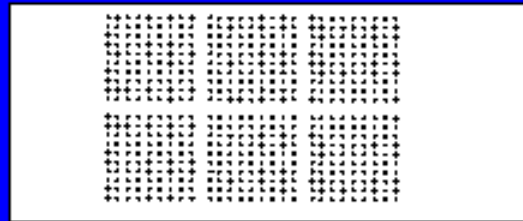
Cy5 Probe



Control

mRNA

Cy3 Probe

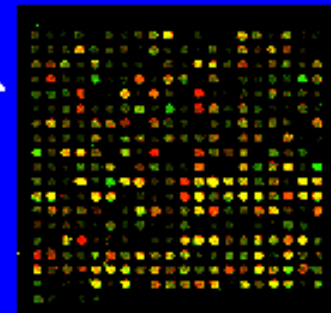


Simultaneous hybridization

Normalization

Data extraction

Scanning



# How to compare 2 cell samples with Two-Color Microarrays?

- ❑ mRNA from sample 1 is extracted and labeled with a **red fluorescent** dye.
- ❑ mRNA from sample 2 is extracted and labeled with a **green fluorescent** dye.
- ❑ Mix the samples and apply it to every spot on the microarray. Hybridize sample mixture to probes.
- ❑ Use optical detector to measure the amount of **green** and **red** fluorescence at each spot.

# Sources of Variations & Experimental Errors

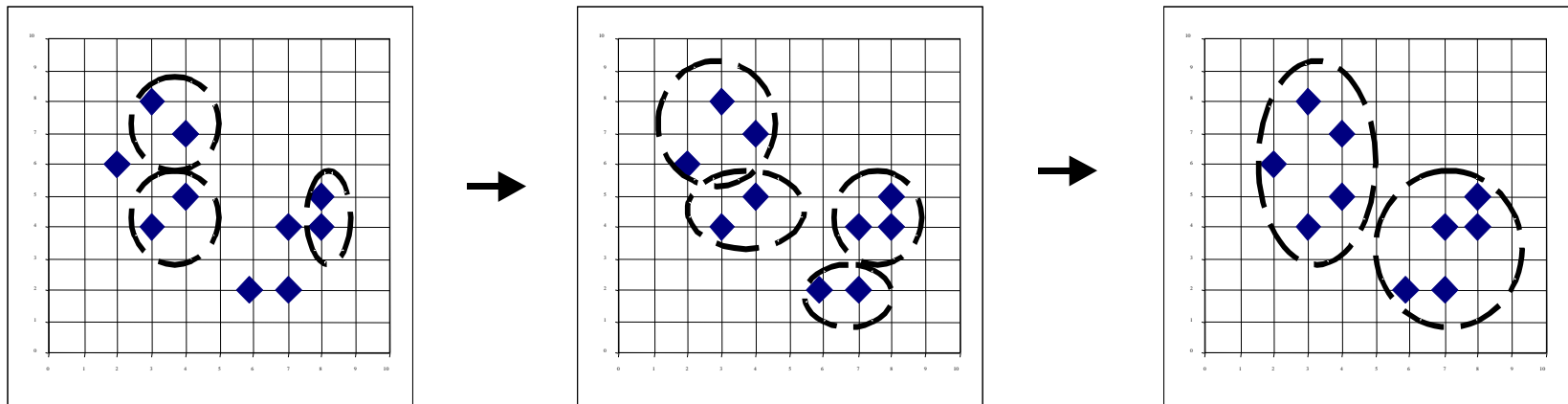
- ❑ Variations in cells/individuals
- ❑ Variations in mRNA extraction, isolation, introduction of dye, variation in dye incorporation, dye interference
- ❑ Variations in probe concentration, probe amounts, substrate surface characteristics
- ❑ Variations in hybridization conditions and kinetics
- ❑ Variations in optical measurements, spot misalignments, discretization effects, noise due to scanner lens and laser irregularities
- ❑ Cross-hybridization of sequences with high sequence identity
- ❑ Limit of factor 2 in precision of results
- ❑ Variation changes with intensity: larger variation at low or high expression levels

Need to Normalize data

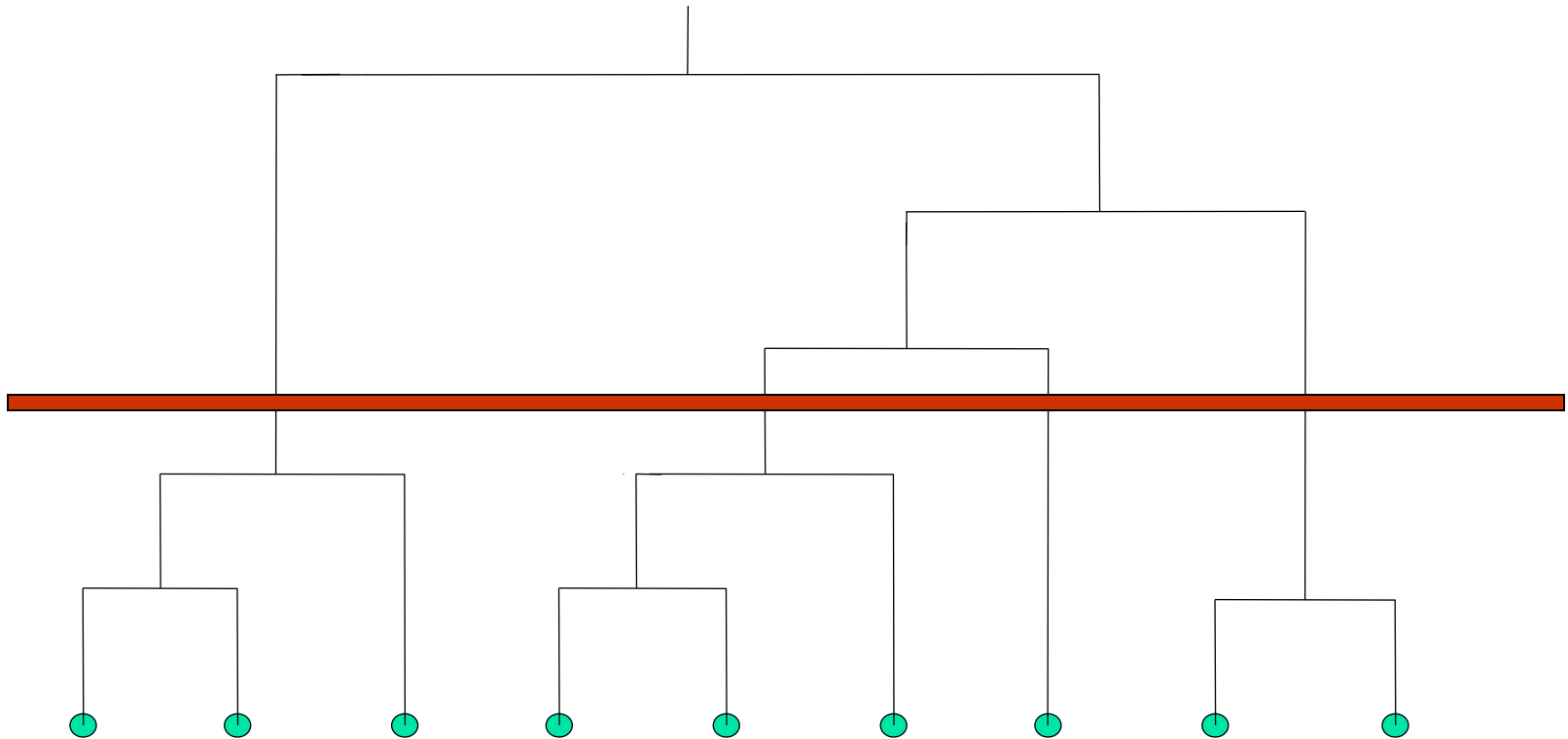
# Clustering

- Clustering is a general method to study patterns in gene expressions.
- Several known methods:
  - *Hierarchical Clustering* (Bottom-Up Approach)
  - *K-means Clustering* (Top-Down Approach)
  - *Self-Organizing Maps (SOM)*

# Hierarchical Clustering: Example



# A Dendrogram



# Hierarchical Clustering [Johnson, SC, 1967]

- Given  $n$  points in  $\mathbb{R}^d$ , compute the distance between every pair of points
- While (not done)
  - Pick closest pair of points  $s_i$  and  $s_j$  and make them part of the same cluster.
  - Replace the pair by an average of the two  $s_{ij}$

Try the applet at:

[http://home.dei.polimi.it/matteucc/Clustering/tutorial\\_html/AppletH.html](http://home.dei.polimi.it/matteucc/Clustering/tutorial_html/AppletH.html)

# Distance Metrics

□ For clustering, define a distance function:

● **Euclidean distance metrics**

$$D_k(X, Y) = \left[ \sum_{i=1}^d (X_i - Y_i)^k \right]^{1/k}$$

k=2: Euclidean Distance

● **Pearson correlation coefficient**

$$\rho_{xy} = \frac{1}{d} \sum_{i=1}^d \left( \frac{X_i - \bar{X}}{\sigma_x} \right) \left( \frac{Y_i - \bar{Y}}{\sigma_y} \right)$$

$-1 \leq \rho_{xy} \leq 1$



**EXHIBIT 3.4** Joint Probability Model for the Ratings of Two People

(a)  $\rho_{XY} = 0$

x	y			Total
	1	2	3	
3	1/9	1/9	1/9	1/3
2	1/9	1/9	1/9	1/3
1	1/9	1/9	1/9	1/3
Total	1/3	1/3	1/3	1

(b)  $\rho_{XY} = \frac{1}{2}$

x	y			Total
	1	2	3	
3	1/18	1/18	4/18	1/3
2	1/18	4/18	1/18	1/3
1	4/18	1/18	1/18	1/3
Total	1/3	1/3	1/3	1

(c)  $\rho_{XY} = -\frac{1}{2}$

x	y			Total
	1	2	3	
3	4/18	1/18	1/18	1/3
2	1/18	4/18	1/18	1/3
1	1/18	1/18	4/18	1/3
Total	1/3	1/3	1/3	1

(d)  $\rho_{XY} = \frac{1}{3}$

x	y			Total
	1	2	3	
3	1/27	2/27	6/27	1/3
2	2/27	5/27	2/27	1/3
1	6/27	2/27	1/27	1/3
Total	1/3	1/3	1/3	1

(e)  $\rho_{XY} = -\frac{2}{3}$

x	y			Total
	1	2	3	
3	6/27	2/27	1/27	1/3
2	2/27	5/27	2/27	1/3
1	1/27	2/27	6/27	1/3
Total	1/3	1/3	1/3	1

(f)  $\rho_{XY} = \frac{2}{3}$

x	y			Total
	1	2	3	
3	1/36	2/36	9/36	1/3
2	2/36	8/36	2/36	1/3
1	9/36	2/36	1/36	1/3
Total	1/3	1/3	1/3	1

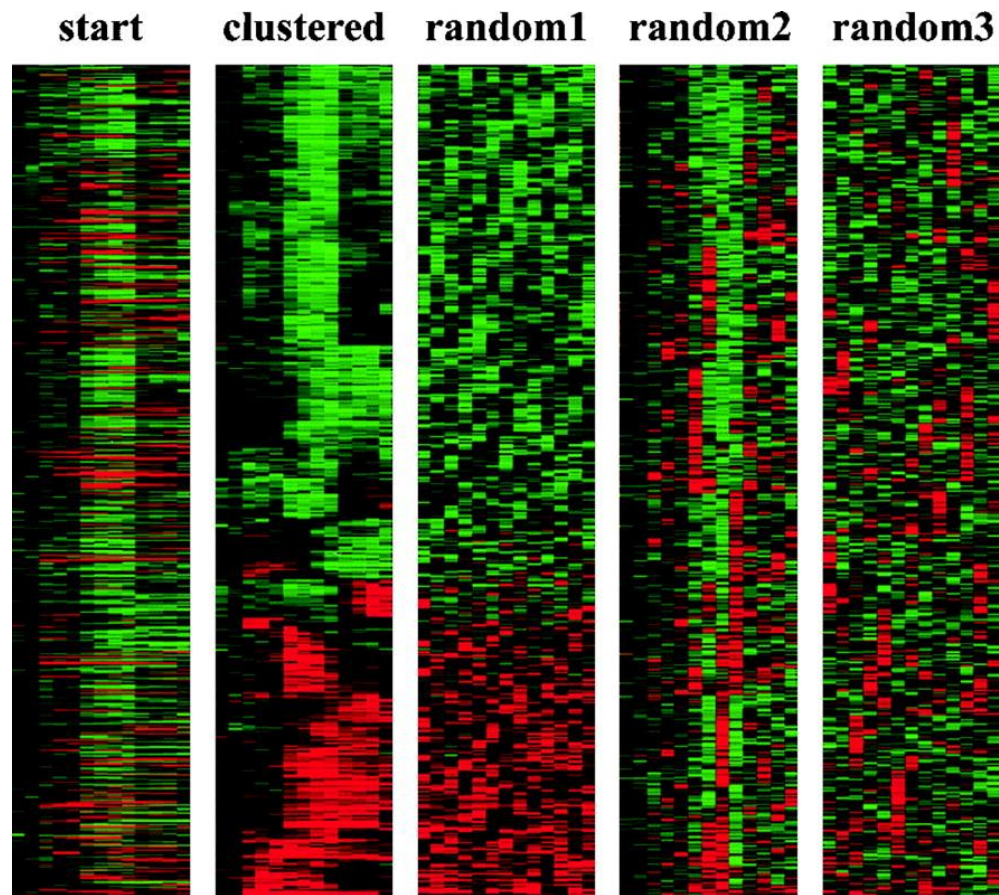
(g)  $\rho_{XY} = -\frac{1}{3}$

x	y			Total
	1	2	3	
3	9/36	2/36	1/36	1/3
2	2/36	8/18	2/18	1/3
1	1/36	2/36	9/36	1/3
Total	1/3	1/3	1/3	1

# Clustering of gene expressions

- Represent each gene as a vector or a point in  $d$ -space where  $d$  is the number of arrays or experiments being analyzed.

# Clustering Random vs. Biological Data



From Eisen MB, et al, PNAS 1998 95(25):14868

# K-Means Clustering: Example

Example from Andrew Moore's tutorial on Clustering.