

## Transcription-regulation (2)

#### Resources



This lectureCooper, pp. 124--132

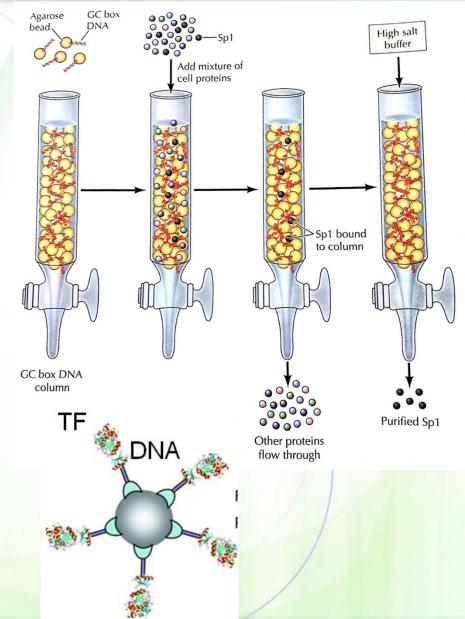
### How do different cells appear?

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- All cells in one body have the same genome, but cell types in a multicellular organism become different from one another because they synthesize different sets of RNA and protein molecules.
- The patterns of mRNA can differentiate cell types from each other.
- How can we investigate RNA expression and function?

# Isolation of transcription factors using affinity chromatography

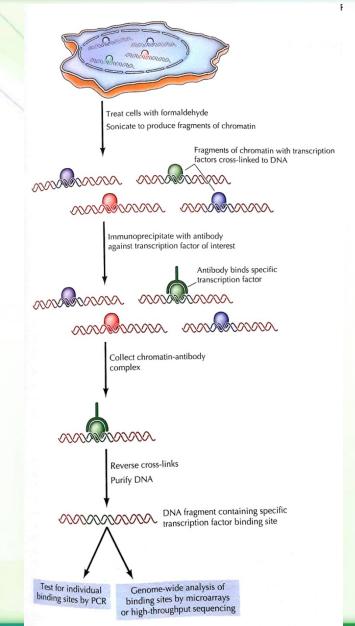
- Attach a DNA fragment with a specific sequence to a bead.
- Pass through nuclear fraction of a cell.
- Elute the bound proteins.



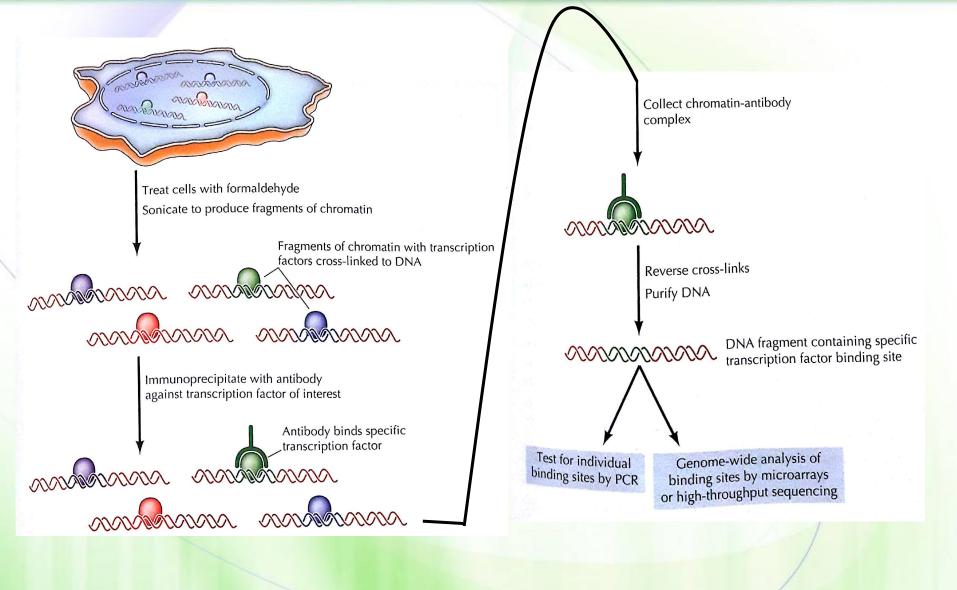
## **Chromatin immunoprecipitation**



- Purpose: identify the DNA sequence to which a transcription factor binds.
- Crosslink proteins bound to DNA.
- Fragments the DNA into smaller pieces.
- Isolate a specific transcription factor by immunoprecipitation.
- Sequence the purified DNA fragment or identify by DNA microarray.



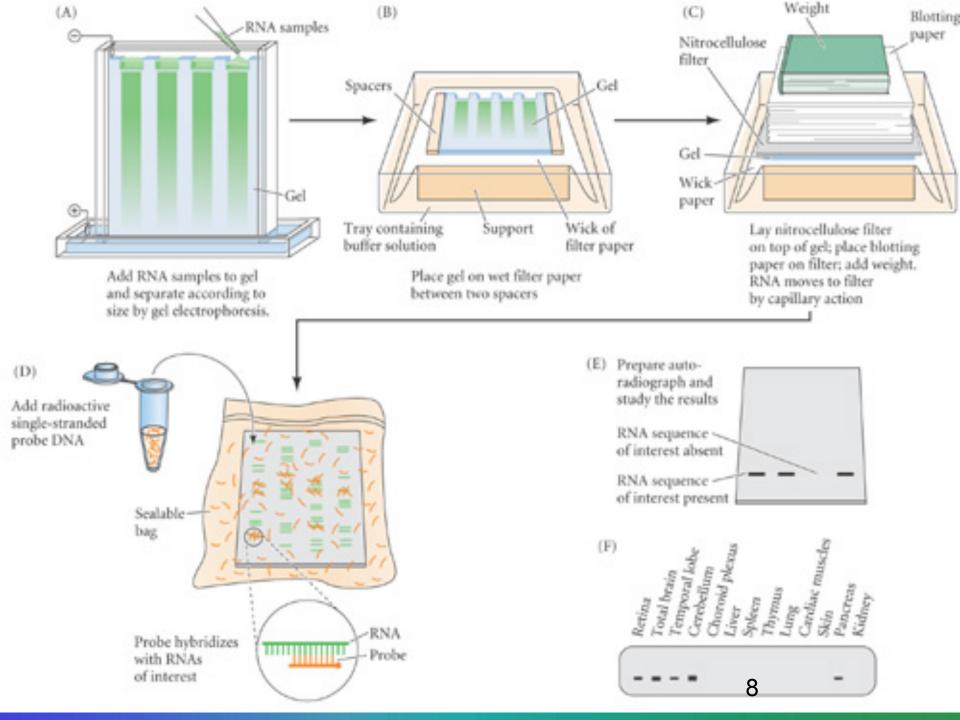




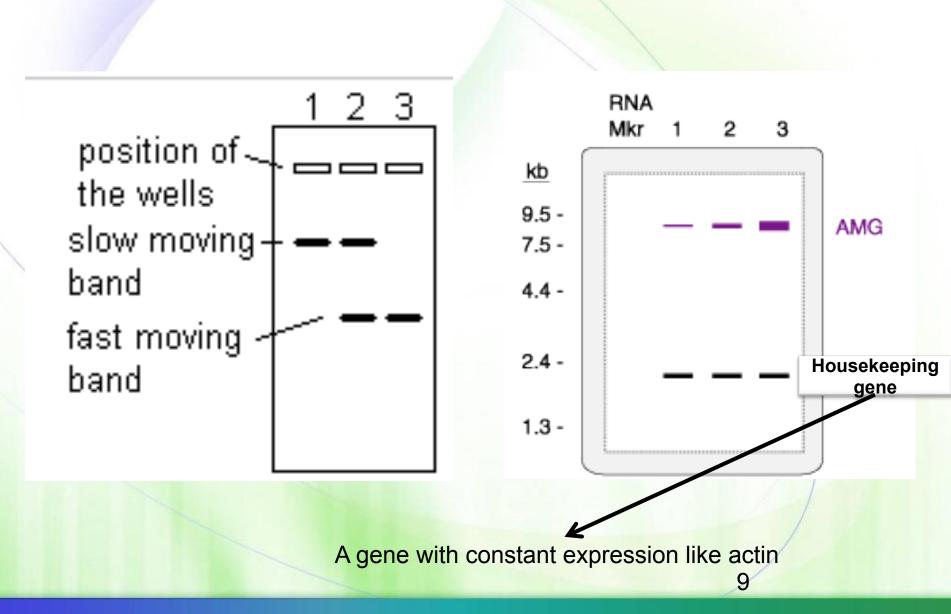
### **Northern blotting**



- This is done exactly like Southern blotting except that RNA from cells is isolated instead of DNA.
- Then RNA molecules are fractionated based on sizes by gel electrophoresis.
- The fractionated RNA molecules are transferred to a membrane.
- The RNA molecules on the membrane are targeted by a labeled DNA probe whose sequence is complementary to a specific RNA molecule.





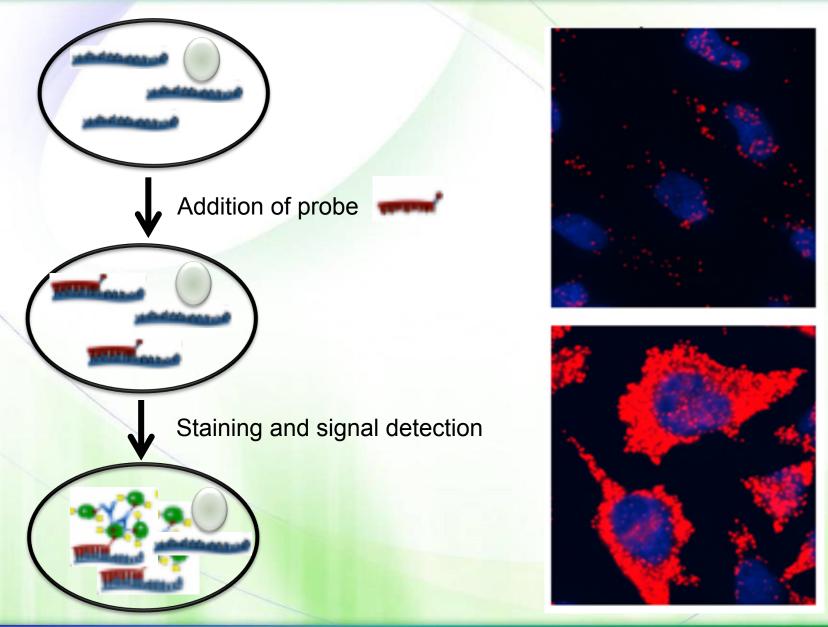


In situ hybridization methods reveals the distribution of specific RNA molecules in cells in tissues.

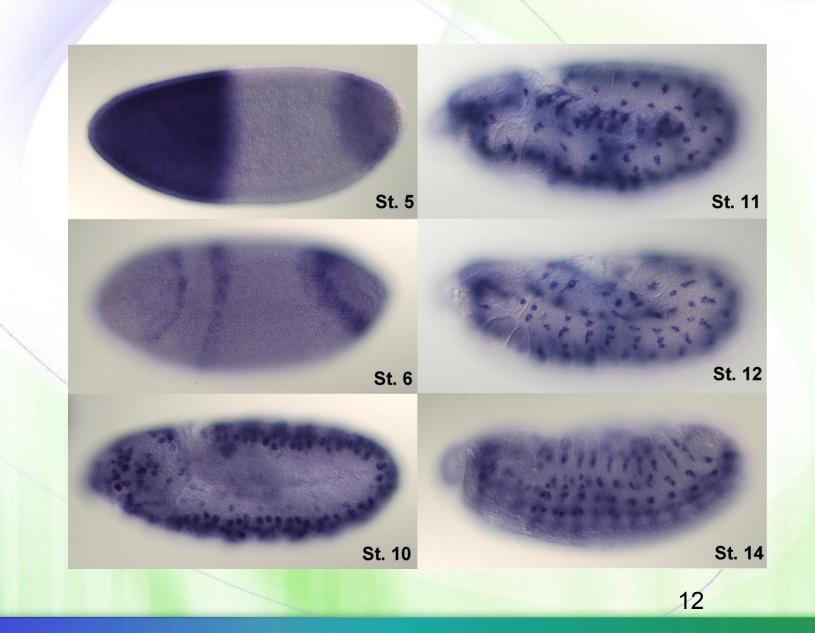
- RNA molecules can hybridize when the tissue is incubated with a complementary DNA or RNA probe.
- In this way the patterns of differential gene expression can be observed in tissues, and the location of specific RNAs can be determined in cells.

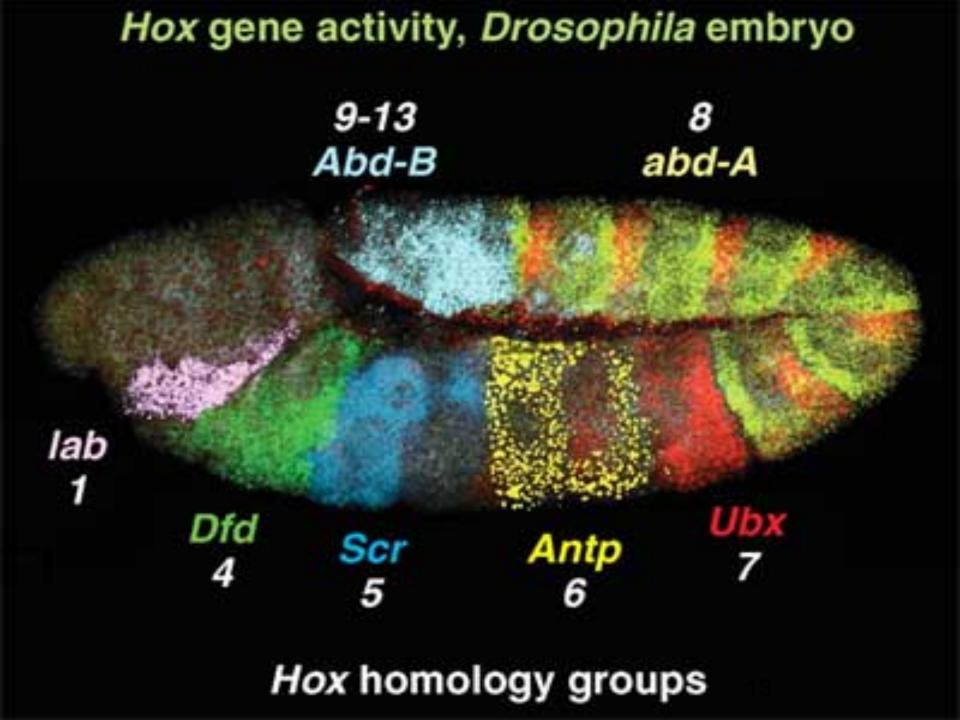
### **Procedure of in situ hybridization**





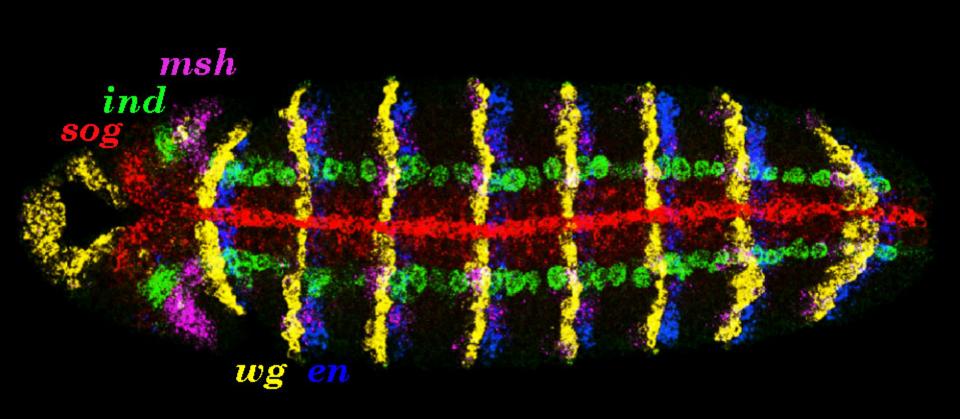
#### **RNA expression in drosophila**







#### Multiplex detection of 5 different transcripts in a single embryo

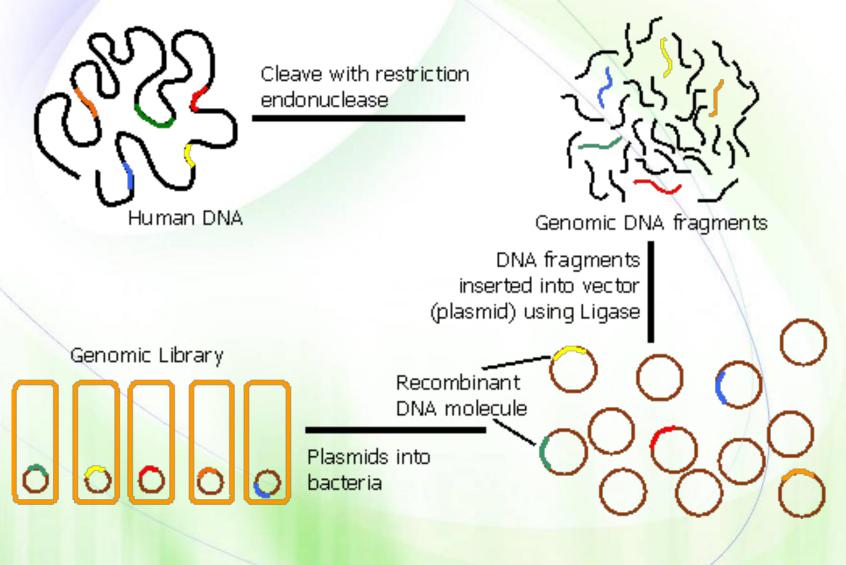


#### What is a DNA library?

- A library can be created for DNA fragments just like book libraries.
- You can have clones of bacteria each containing a specific piece of DNA.
- You can save these clones in the freezer and take whichever clone you want to study.
- <u>http://www.sumanasinc.com/webcontent/animations/</u> <u>content/dnalibrary.html</u>

#### **Genomic DNA library**

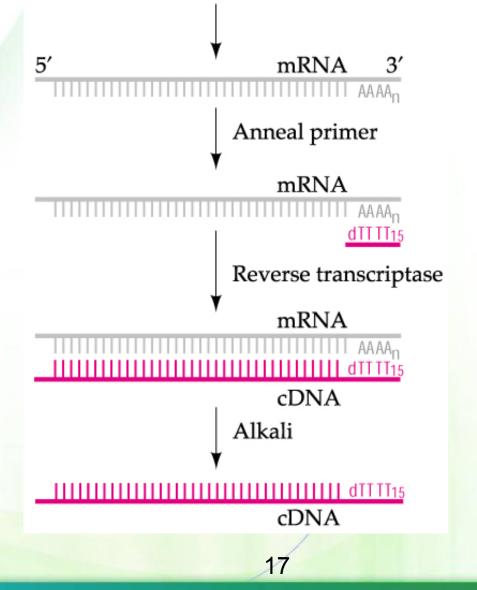




#### **cDNA** library



- This library contains only those DNA sequences that are transcribed into mRNA.
- This is done by extracting the mRNA from cells and then making a complementary DNA (cDNA) copy of each mRNA molecule present.
- cDNA is made by retroviral reverse transcriptase.



#### **Genomic vs. cDNA libraries**

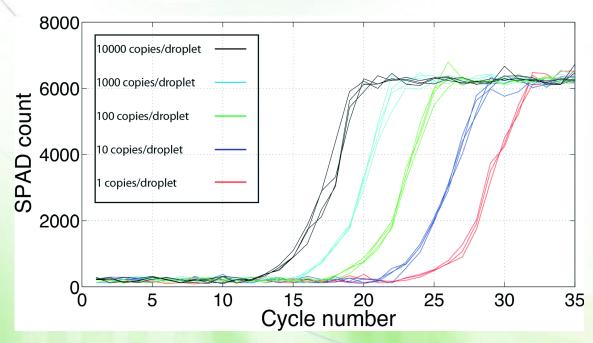


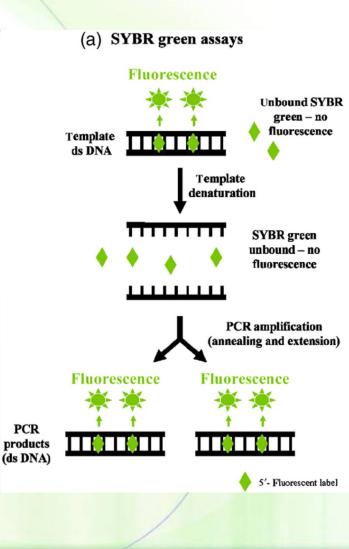
- Genomic clones represent a random sample of all of the DNA sequences in an organism. By contrast, cDNA clones contain only those regions of the genome that have been transcribed into mRNA.
  - Because the cells of different tissues produce distinct sets of mRNA molecules, a distinct cDNA library is obtained for each type of cell used to prepare the library.

#### **Quantitative real-time PCR of mRNA**



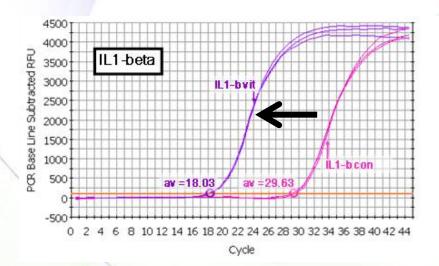
- Another way of relative quantitation of RNA expression is by converting RNA into cDNA followed by PCR in the presence of SYBR green.
- The higher the amount of RNA (cDNA), the sooner it is detected.



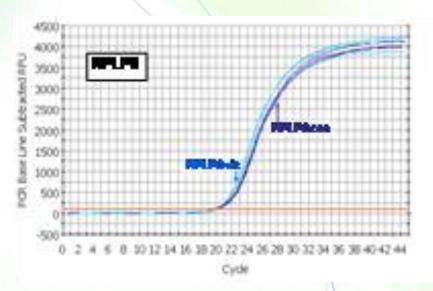




#### A gene of interest



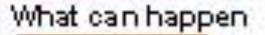
#### Housekeeping gene

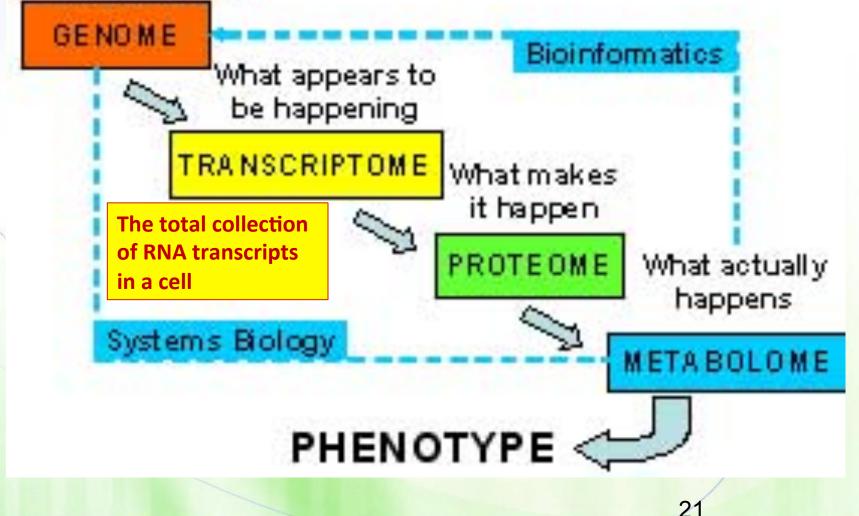


#### **Unaltered expression**

#### **The science of -omics**







#### **Studying the transcriptome**



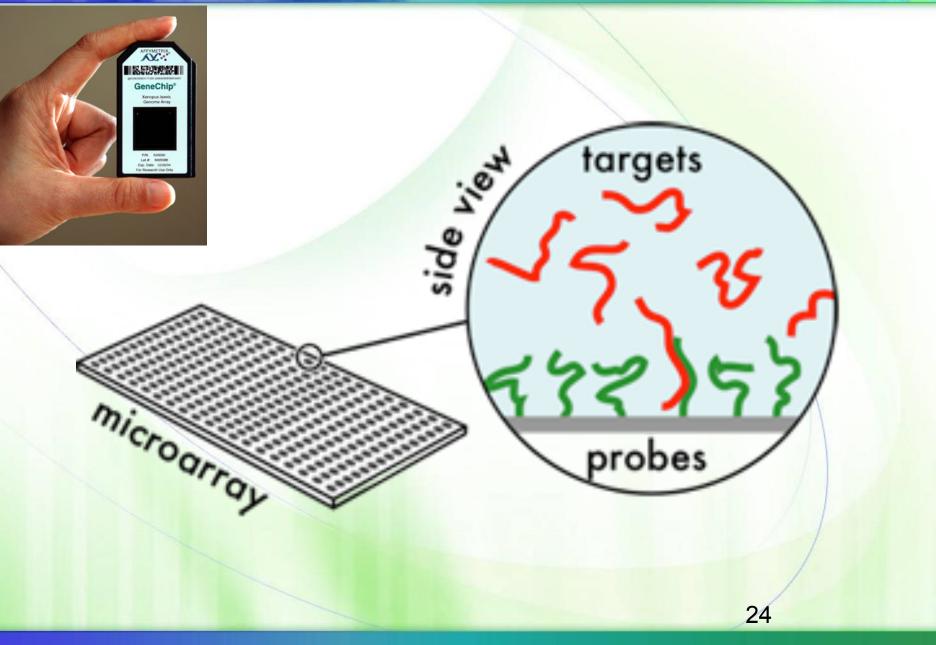
- One such method in studying transcriptomes is DNA microarrays, which allow the analysis of the RNA products of thousands of genes all at once.
  - By examining the expression of so many genes simultaneously, we can understand gene expression patterns in physiological and pathological states.

#### **DNA microarrays**



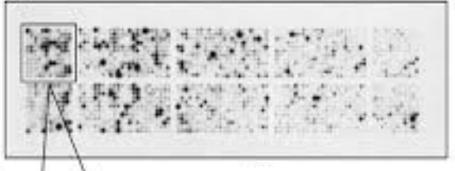
- DNA microarrays are glass microscope slides spotted with up to tens of thousands of DNA fragments in an area the size of a fingernail.
- The exact sequence and position of every DNA fragment on the array is known.
- http://learn.genetics.utah.edu/content/labs/ microarray/
- <u>http://www.sumanasinc.com/webcontent/animations/</u> <u>content/dnachips.html</u>





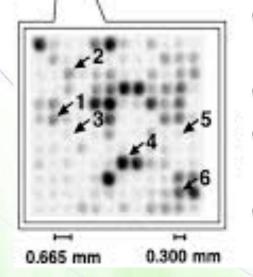
### A DNA microarray





This is done for a single sample using radioactively labeled cDNA.

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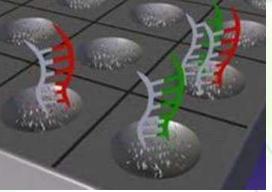


- mRNA from the cells being studied is first extracted and converted to cDNA.
- The cDNA is labeled with a radioactive probe.
- The microarray is incubated with the labeled cDNA sample for hybridization to occur.
- If a gene is expressed, then the cDNA will exist and bind to a specific complementary DNA fragment on the microarray.
- Binding can be detected since the cDNA is labeled and expression is determined.

#### **Comparative expression**



- In order to compare expression of genes two different samples, the cDNA molecules are fluorescently labeled with different colors (green and red) and added to the array.
- An increases in the amount of a RNA molecule in one sample versus the other is reflected by an increase the amount of produced cDNA and an increase in fluorescence in the bound spot.

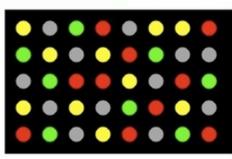


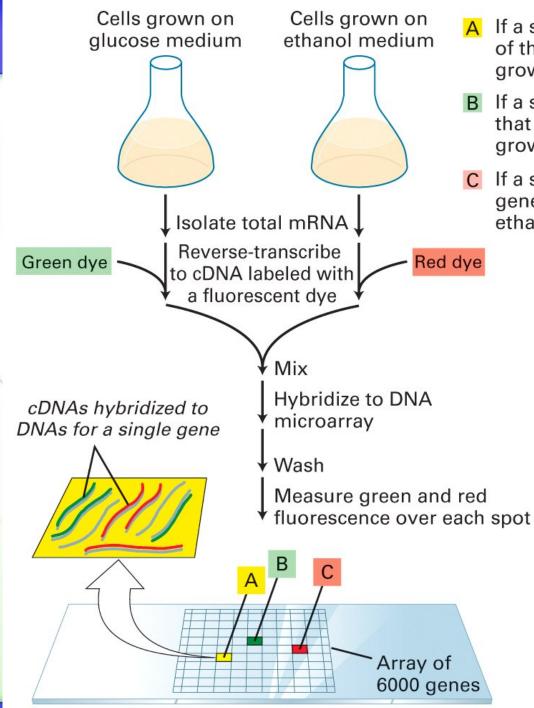
- Gene NOT active in either normal or diseased sample
- Gene IS active in both normal and diseased sample



- Gene active in normal only  $\Rightarrow$  very interesting!
- Gene active in disease only ⇒ very interesting!

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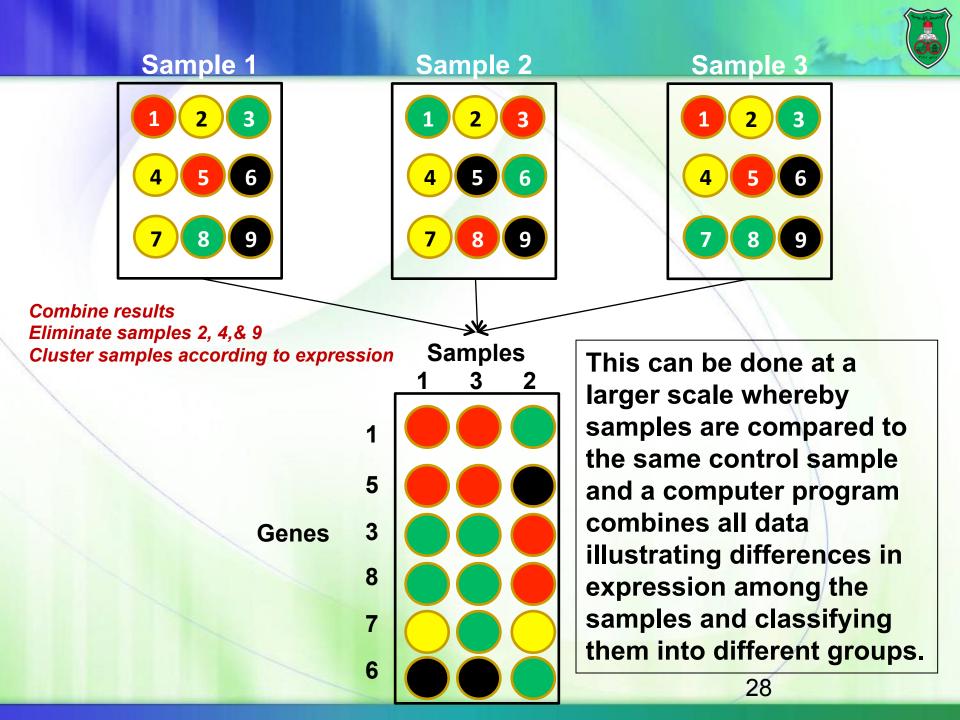


A If a spot is yellow, expression of that gene is the same in cells grown either on glucose or ethanol



- B If a spot is green, expression of that gene is greater in cells grown in glucose
- C If a spot is red, expression of that gene is greater in cells grown in ethanol

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#### **DNA microarrays and breast cancer**

