

## Basic lab techniques

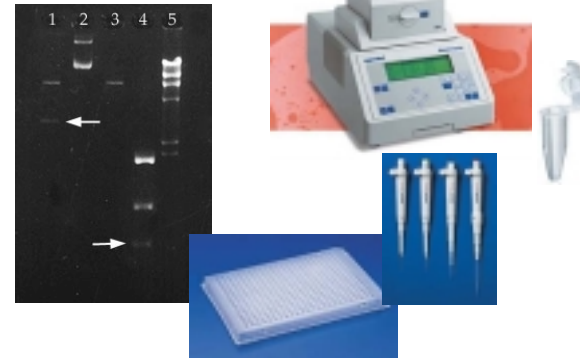
Sandrine Dudoit

Bioconductor short course  
Summer 2002



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## Lab techniques



## Basic lab techniques for nucleic acids

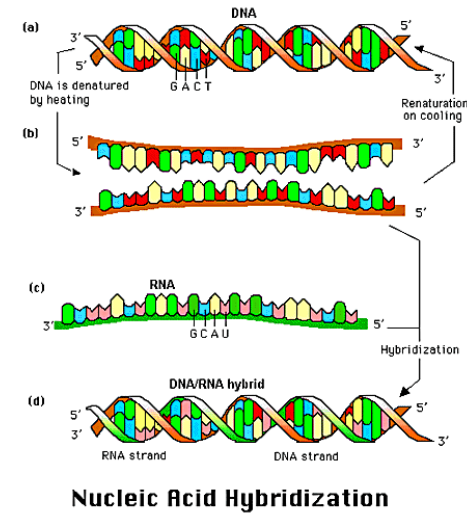
- Hybridization.
- Cut: restriction enzymes.
- Amplify: PCR.
- Sort: gel electrophoresis.
- Probe: blots and microarrays.

## Why?

- Why cut, amplify, sort, probe?
  - Sequencing;
  - Genotyping (cf. genetic mapping, forensics);
  - Measuring gene expression;
  - Etc.

## Hybridization

- **Hybridization** refers to the **annealing** of two nucleic acid strands following the base-pairing rules.
- Nucleic acid strands in a duplex can be separated, or **denatured**, by heating to destroy the hydrogen bonds.



## Restriction enzymes

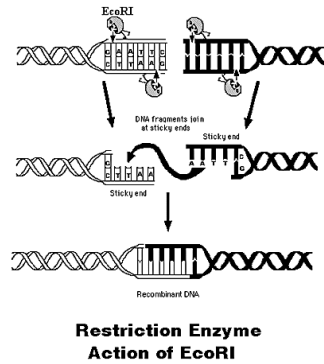
- DNA **restriction enzymes** or **restriction endonucleases** recognize short, specific sequences of DNA bases and make breaks in the sugar-phosphate backbone of the DNA.
- The recognition sites are usually **palindromes**, .i.e, the sequence in one strand is the same as that in the other strand, read in the reverse direction.
- Some restriction enzymes make staggered cuts in the opposite strand, creating complementary, single-stranded ends or **sticky ends**; others cut across both strands creating DNA fragments with **blunt ends**.

## EcoRI

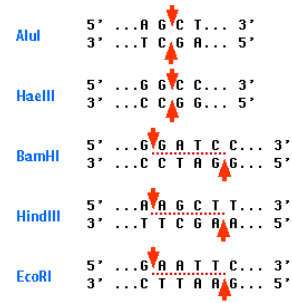
- Restriction enzymes allow bacteria to self-defend against invading DNA-containing organisms (e.g. virus).
- EcoRI, from *Escherichia coli* or *E. coli*.

5' G|AATTC  
3' CTTAA|G

## Restriction enzymes



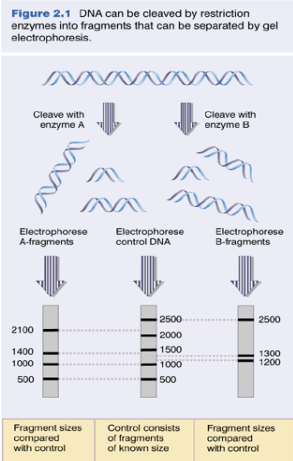
## Restriction enzymes



**AluI** and **HaeIII** produce blunt ends

**BamHI** **HindIII** and **EcoRI** produce "sticky" ends <http://www.ultranet.com/~jkimball/BiologyPages/>

## Restriction enzymes



## PCR

- **Polymerase chain reaction** or **PCR** is a widely used technique for creating billions of copies, i.e., **amplifying**, a single DNA fragment.
- It is based on nucleic acid hybridization.

## PCR

- PCR relies on
  - Known sequence for the 3' end of the **template**, i.e., segment to be amplified.
  - Availability of **primers**, i.e., synthetic oligonucleotides complementary to the 3' ends of the template.
  - Use of **temperature** to control DNA **annealing** and **denaturation**.
  - Existence of a temperature resistant enzyme for DNA synthesis by primer extension: **Taq polymerase** (*Thermus aquaticus*, bacterium found in Yellowstone hot springs).

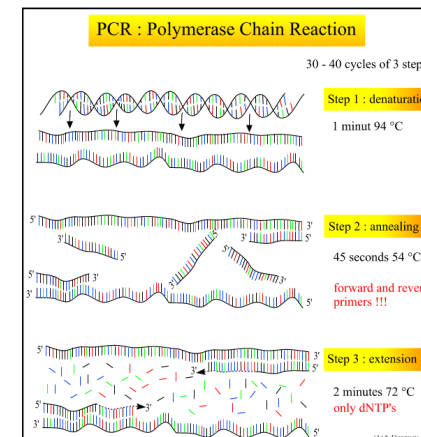
## PCR

- Main ingredients:
  - DNA template,
  - primers in great excess of template,
  - dNTPs: deoxynucleotide triphosphates,
  - Taq polymerase.
- Repeated **cycles** of DNA denaturation (heating) and synthesis (cooling) rapidly provide many copies of the template.
- There are three major steps in a PCR, which are repeated for 30 or 40 cycles.

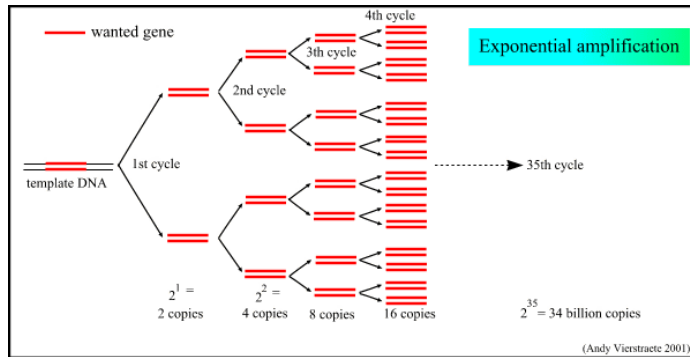
## PCR

1. **Denaturation** (94°C): double strand melts open to single-stranded DNA, enzymatic reactions stop.
2. **Annealing** (54°C): Hydrogen bonds form between the single-stranded primer and template, the polymerase attaches to the duplex and starts copying the template.
3. **Extension** (72°C): At the ideal temperature for the polymerase, bases complementary to the template are coupled to the primer on the 3' end (the polymerase adds dNTPs from 5' to 3').

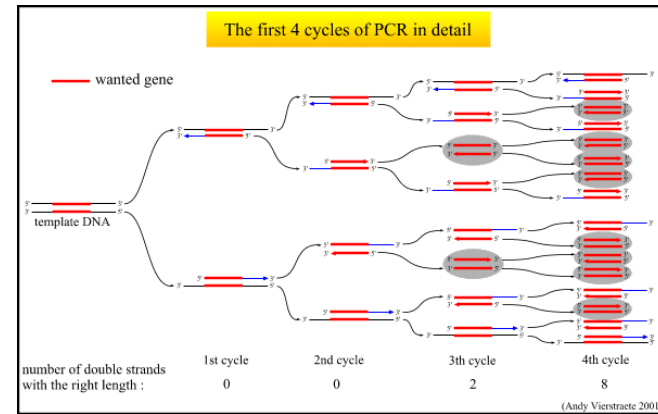
## PCR



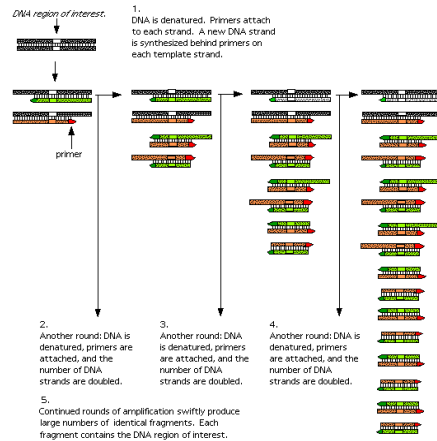
# PCR



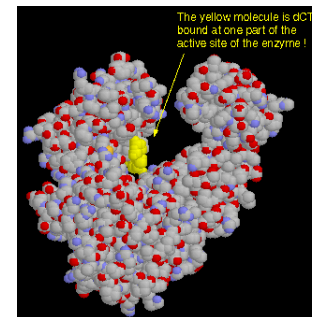
# PCR



# PCR



# Taq polymerase



<http://berget.mcs.cmu.edu/education/TechTeach/replication/TaqL.html>

## Reverse transcriptase PCR

- Amplify **RNA** into DNA.
- E.g. complementary DNA or **cDNA** from mRNA.
- Based on an RNA-dependent DNA polymerase, **reverse transcriptase**, that catalyzes the synthesis of DNA from dNTPs, using RNA as a template.
- The reverse transcriptase enzyme is found in **retroviruses** and is responsible for their replication.

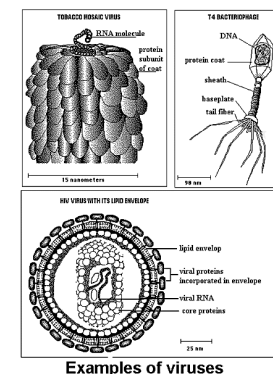
## Viruses and retroviruses

- **Viruses** consist of a **nucleic acid** surrounded by a **protein capsid**.
- **Retroviruses** contain **RNA** as the hereditary material in place of the more common DNA.
- E.g. Human immunodeficiency virus, HIV, the virus that causes AIDS.

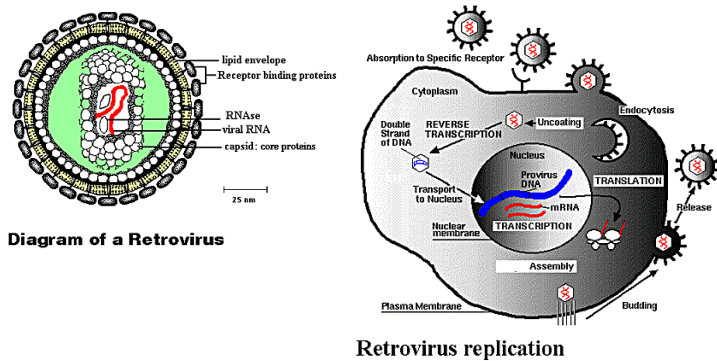
## Retroviruses

- Retroviruses contain the enzyme **reverse transcriptase** (ribonuclease or RNase), which causes synthesis of a complementary DNA molecule (cDNA) using virus RNA as a template.
- When a retrovirus infects a cell, it injects its RNA into the cytoplasm of that cell along with the reverse transcriptase.
- The cDNA produced from the RNA template contains the virally derived genetic instructions and allows infection of the host cell to proceed.

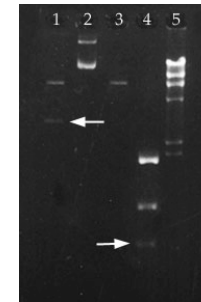
## Viruses



## Retroviruses



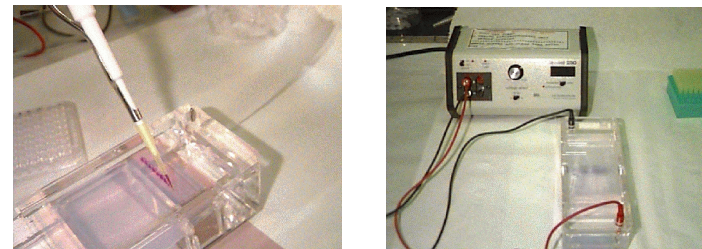
## Gel electrophoresis



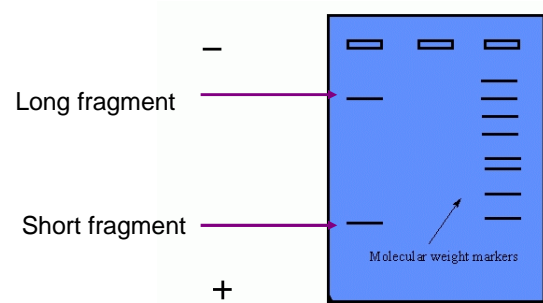
## Gel electrophoresis

- **Electro** refers to electrical field; **phoresis**, from the Greek phoros, means "to carry across".
- **Gel electrophoresis** is a procedure for separating a mixture of charged molecules through a stationary material (gel) in an electrical field.
- Molecules are separated according to electric charge, size, and other physical properties.
- The gel is a colloid in a solid form (e.g. agarose, colloid from seaweed).
- Activated electrodes at either end of the gel provide the driving force.

## Gel electrophoresis



## Gel electrophoresis



## Gel electrophoresis



<http://web.utk.edu/~khughes/>

## Probing

- Goal. Monitor the presence or abundance of specific DNA/RNA sequences in a pool of DNA/RNA (e.g. DNA from a certain type of cells).
- A **probe** is a labeled (radioactive or fluorescent) single-stranded oligonucleotide, synthesized to be complementary to the sequence of interest – i.e., the probe sequence is known.
- The DNA/RNA sample interrogated by the probe is called the **target**.

## Probing

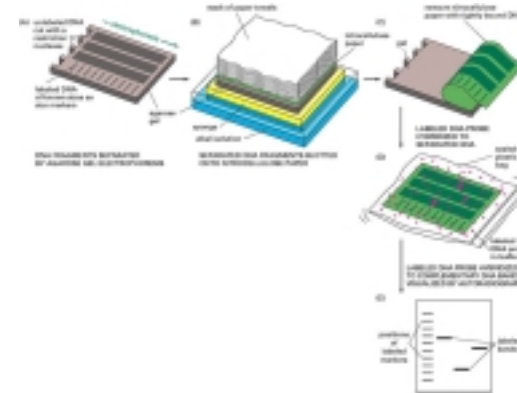
- The probe is attached to a solid support (e.g. membrane) and incubated with the target to allow hybridization of the target to the probe.
- The extent of hybridization of the target to the probe reflects the abundance of the probe in the target.
- Quantification can be done by, e.g., X-ray for radioactive probes.



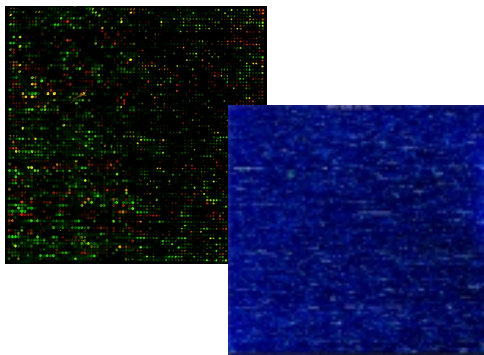
## Blots

- Blots are named for the **target** molecule.
- **Southern blot**: **DNA** cut with restriction enzymes - probed with radioactive DNA.
- **Northern blot**: **RNA** - probed with radioactive DNA or RNA.
- **Western blot**: **protein** - probed with radioactive or enzymatically-tagged antibodies.

## Southern blot



## Microarrays ... blots on a genomic scale



## WWW resources

- **Access Excellence**  
<http://www.accessexcellence.com/AB/GG/>
- **Genes VII**  
<http://www.oup.co.uk/best/textbooks/biochemistry/genesvii/>
- **Human Genome Project Education Resources**  
<http://www.ornl.gov/hgmis/education/education.html>
- **Kimball's Biology Pages**  
<http://www.ultranet.com/~jkimball/BiologyPages/>
- **MIT Biology Hypertextbook**  
<http://esq-www.mit.edu:8001/>
- **PCR**  
<http://www.highveld.com/pcr.html>