Techniques

- Biotechnology: use of microbes to make a protein product
- Recombinant DNA Technology:
  - Insertion or modification of genes to produce desired proteins
- Genetic engineering: manipulation of genes/insert DNA into cells
- Gene Cloning: isolating genes from one organism, manipulating purified DNA in vitro, and transferring to another organism

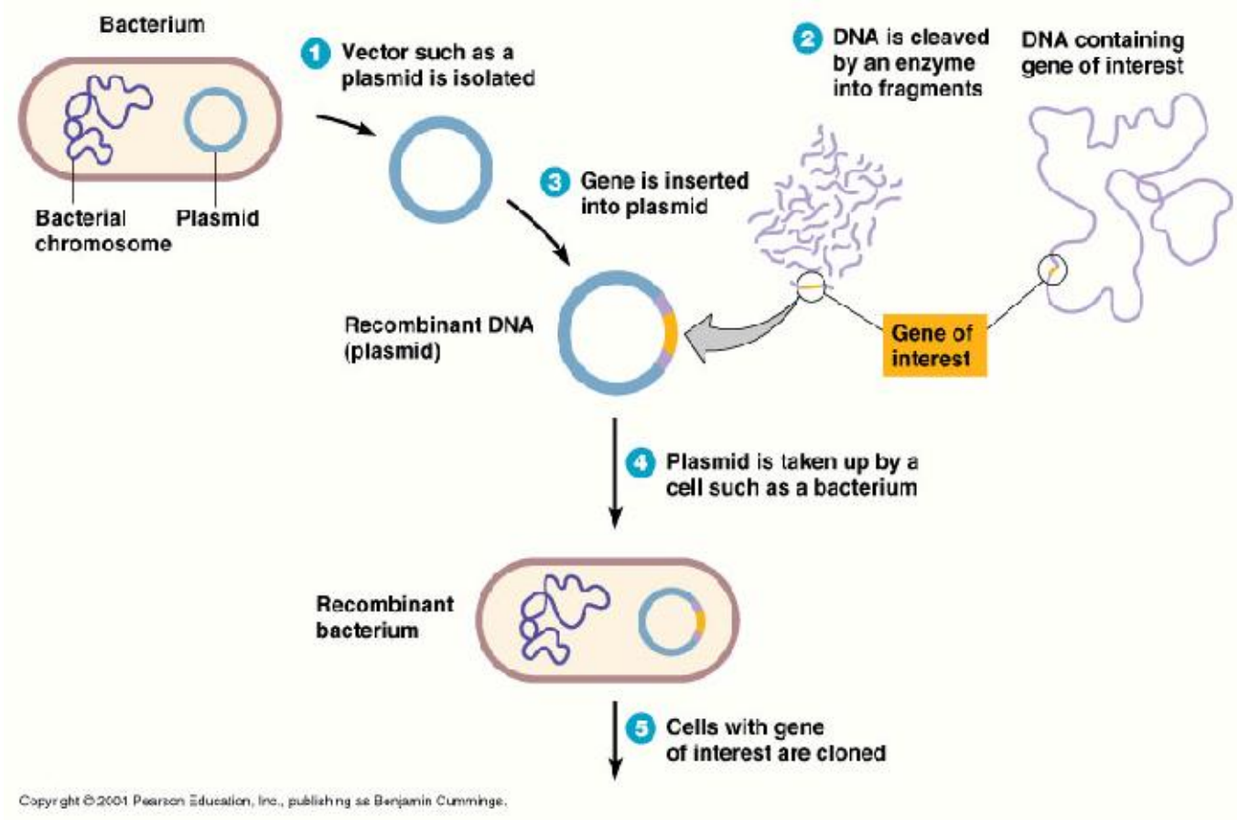
Why is genetic engineering important?

- Purify protein
  - Insulin
  - Growth factor
  - Interferon
- Generate more copies of a particular gene: “amplify DNA”
- Research gene function and regulation

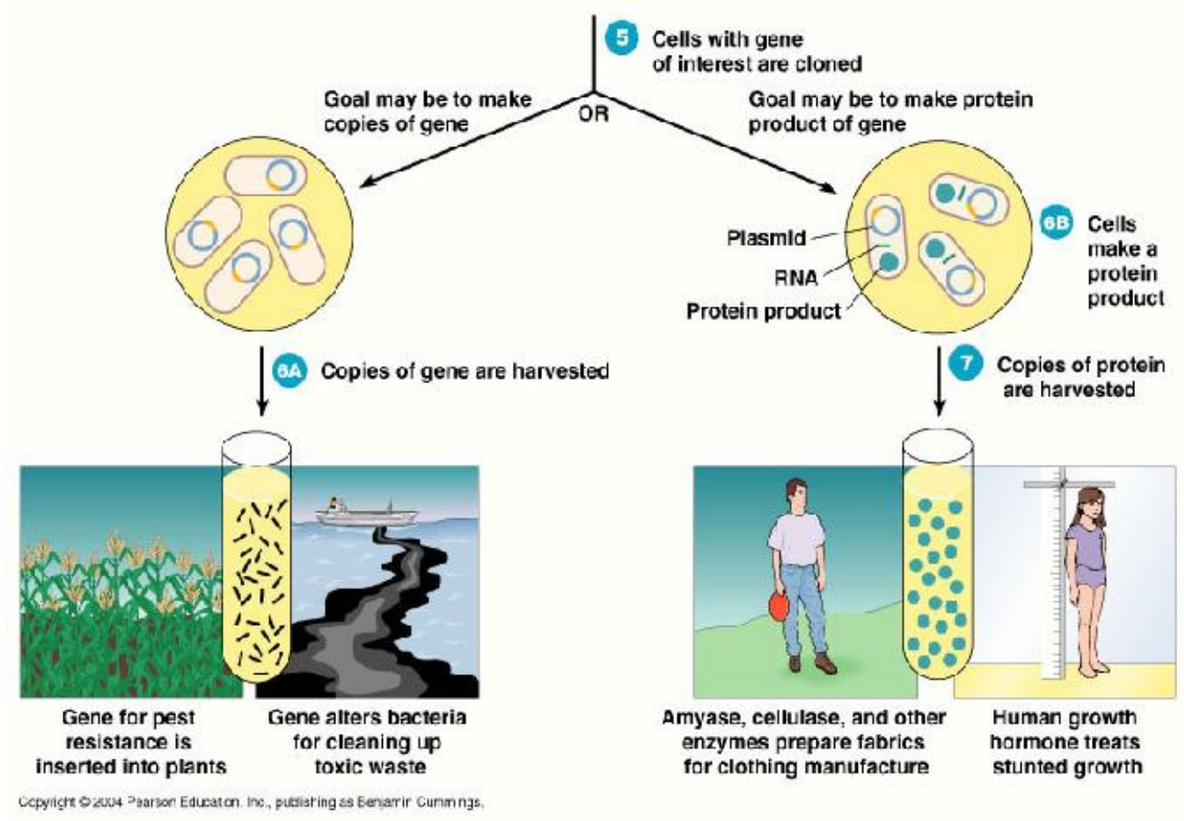
Selection

- Artificial Selection: select breeds or strains with desirable traits (eg. Antibiotic producers)
- Mutation: Mutagens cause mutations that might result in a microbe with a desirable trait
- Site-directed mutagenesis: make specific changes in gene (mutate gene so that an organism can produce more penicillin;
- 1000x more)
- Select and culture microbe with the desired mutation

The process of genetic engineering



The outcomes of genetic engineering



Vectors are types of DNA

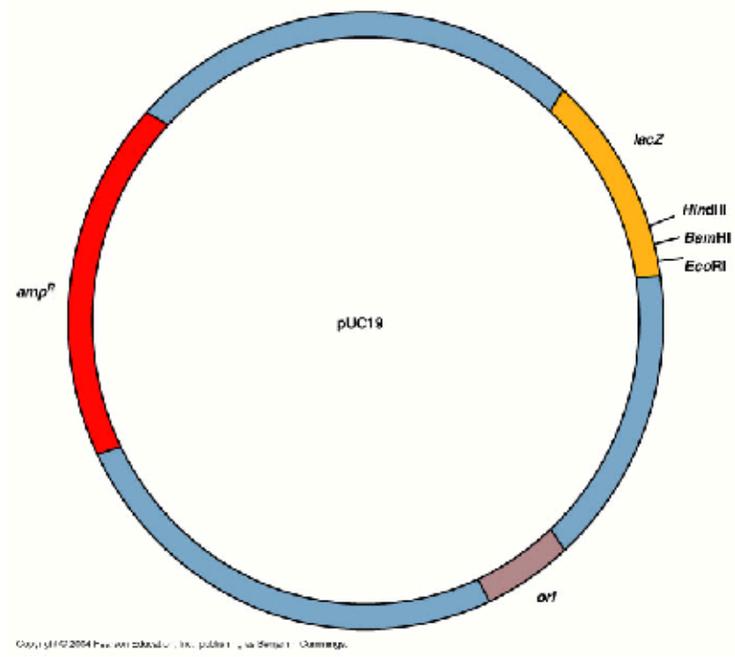
- Must be able to self replicate (WHY?)
- Must contain a promoter region
- Must a reasonable size and circular
- Often contain marker genes (antibiotic resistance genes) for easy identification of cells containing the vector

Types of Vectors

- Viral DNA
  - can carry larger pieces of foreign DNA
- Plasmids
  - pUC19 contains genes for easy selection (lacZ and amp)

Plasmids make good vectors

- pUC19 contains genes for easy selection
- Contains a polylinker region for restriction enzymes
- What happens when the plasmid is cut with EcoR1?

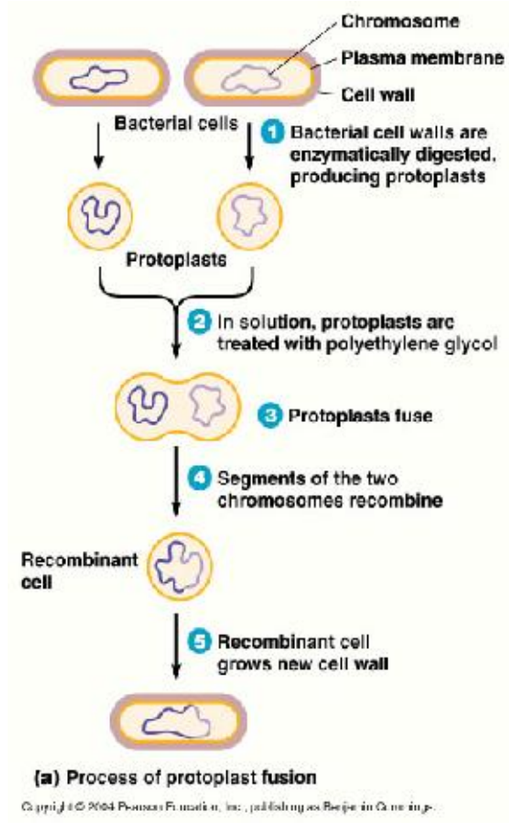


DNA can be inserted into cell by:

- Transformation
  - Naturally competent cells
  - Treat cells (E.coli, yeast, mammal cells) to make competent
    - Soak E.coli in CaCl, mix with DNA, mild heat shock

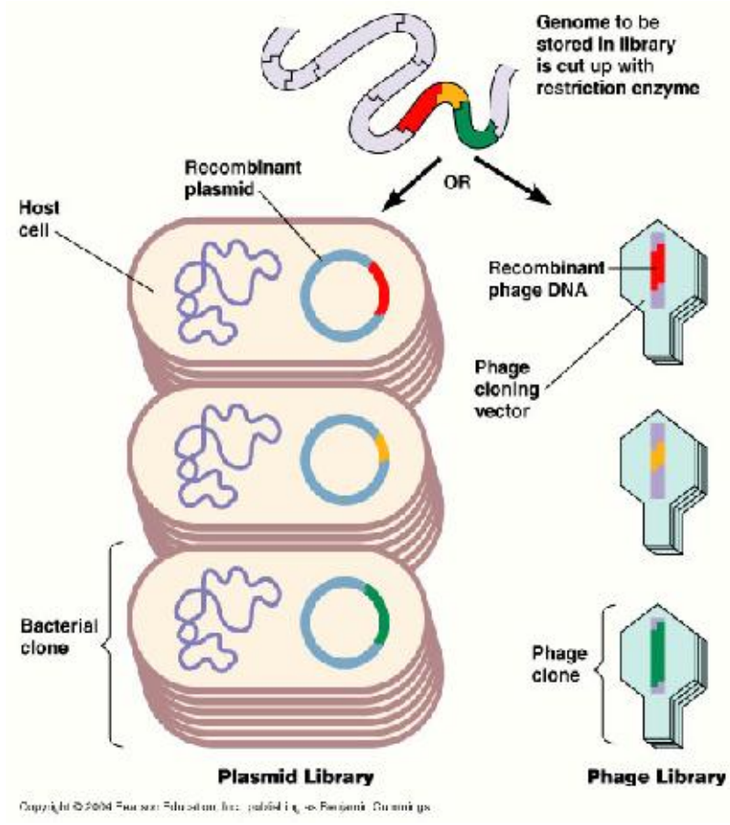
DNA can be inserted into cell by:

- Transformation
- Electroporation
  - Cells with cell wall need to be converted to protoplasts



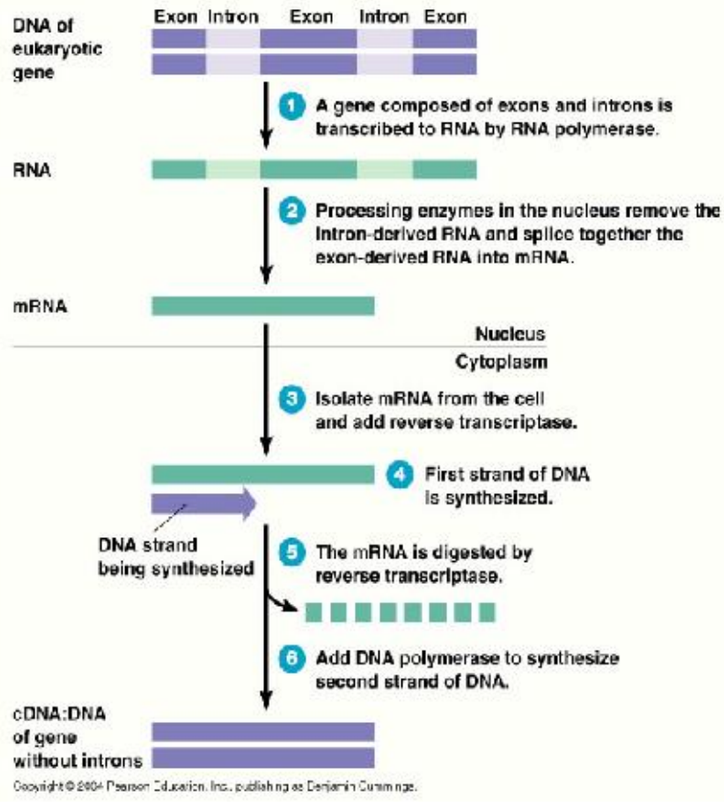
Sources of the DNA that is inserted into the vector?

- Gene library
  - – collection of clones that contain every gene of an organism
  - – pieces of an entire genome stored in plasmids or phages
- Synthesize DNA with a DNA machine



DNA from eukaryotic cells

- cDNA (complementary DNA)
- Problem that genes contain exons and introns
- Use reverse transcriptase synthesize cDNA from mRNA template

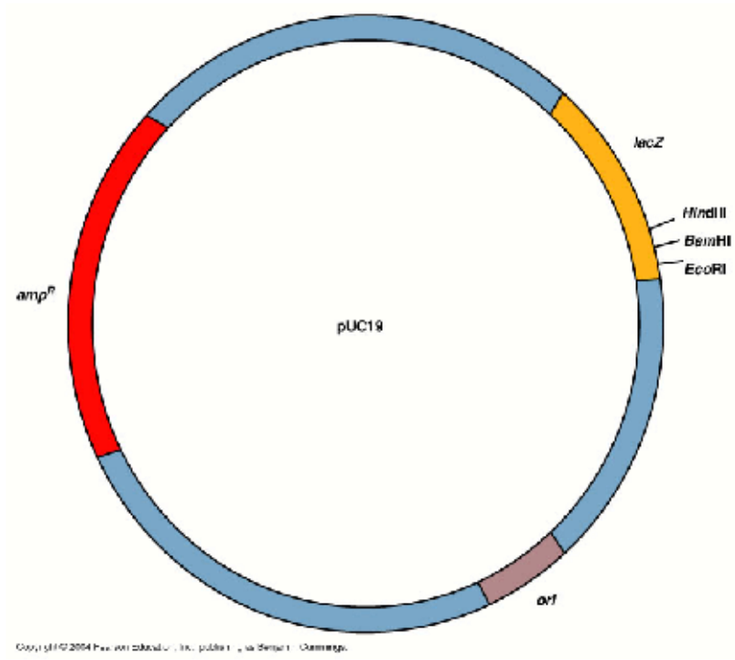


What do we have so far?

- Vectors
- Ways to get DNA into cells
- DNA of "gene of interest"
- Now we need to look at the selection process….how do we find the cells that have taken up the foreign DNA?

A look back at pUC19

- pUC19 has antibiotic resistance gene for anpicillin
- Also has the LacZ gene which codes for Beta-galactosidase
- What happens when cells take up this DNA?



Blue-White Screening

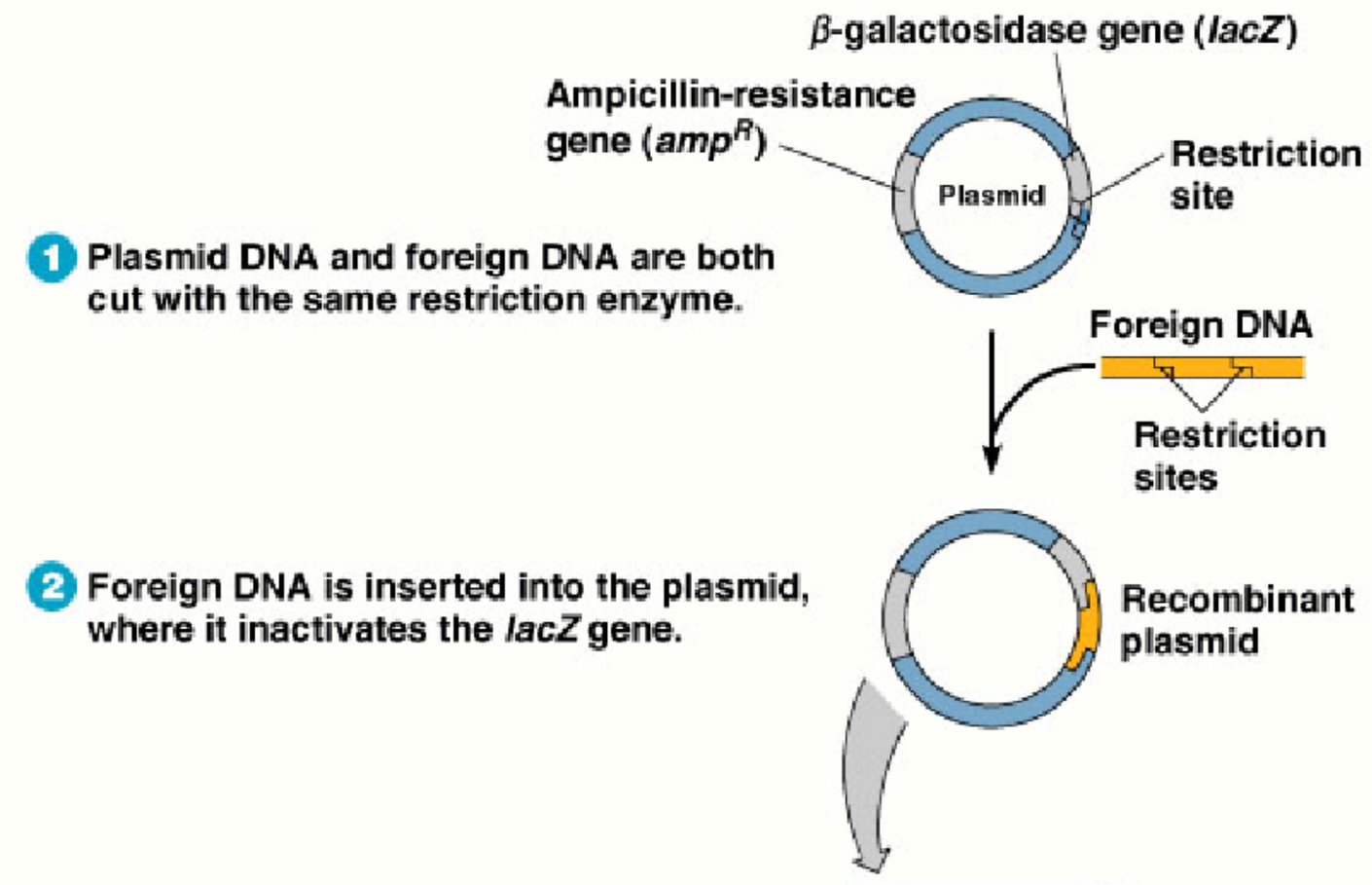


Figure 9.11.1

Blue-White Screening

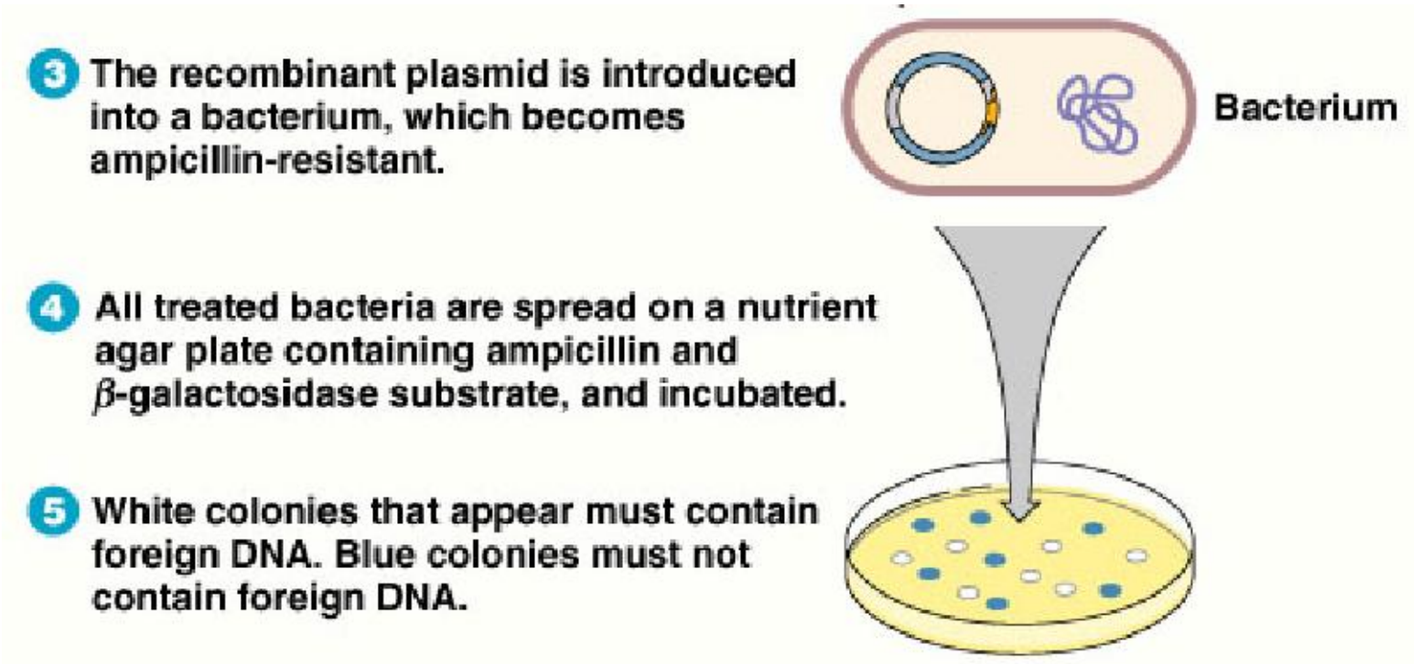
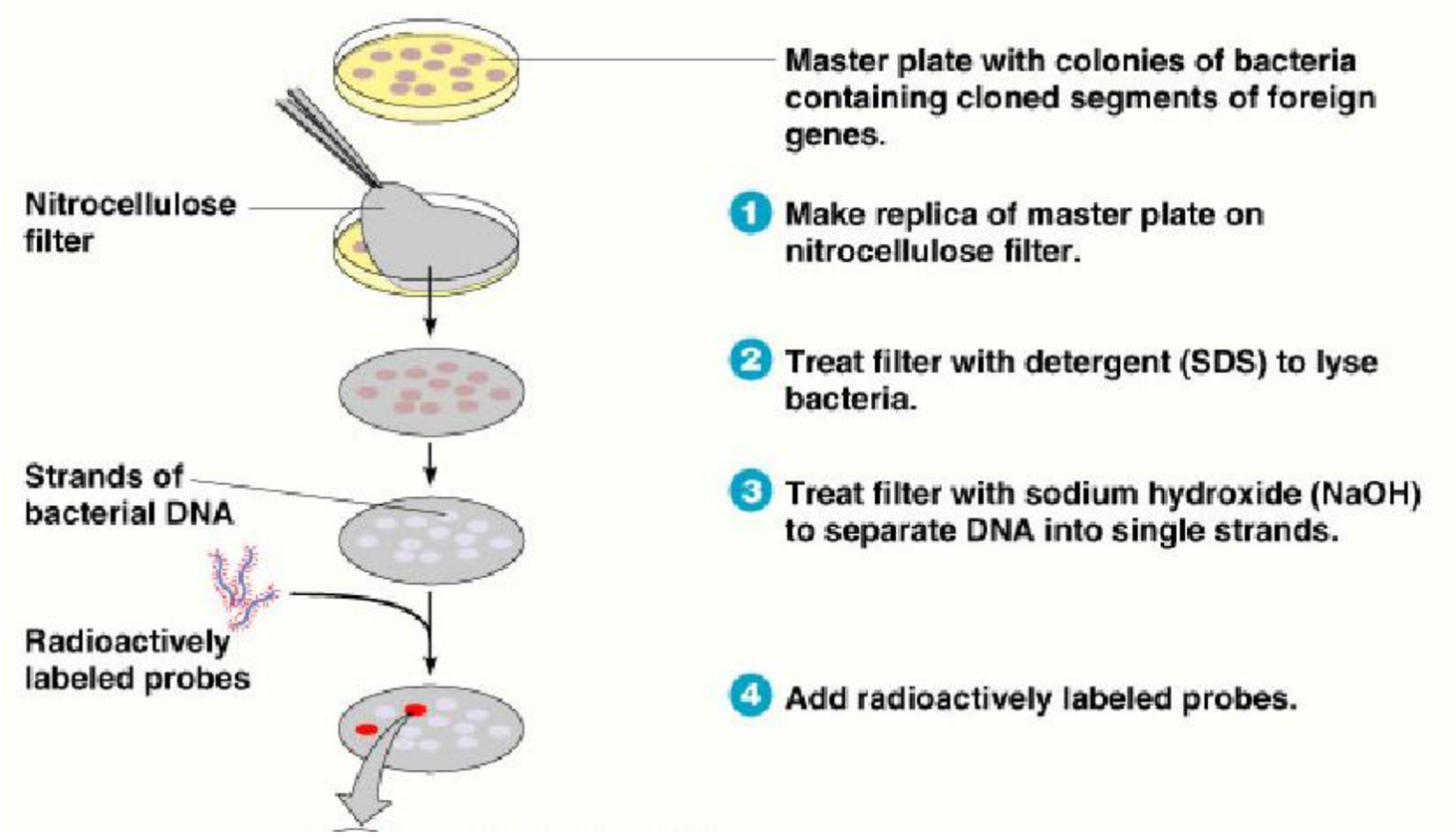


Figure 9.11.2

Colony hybridization for specific gene



Figur

Colony Hybridization

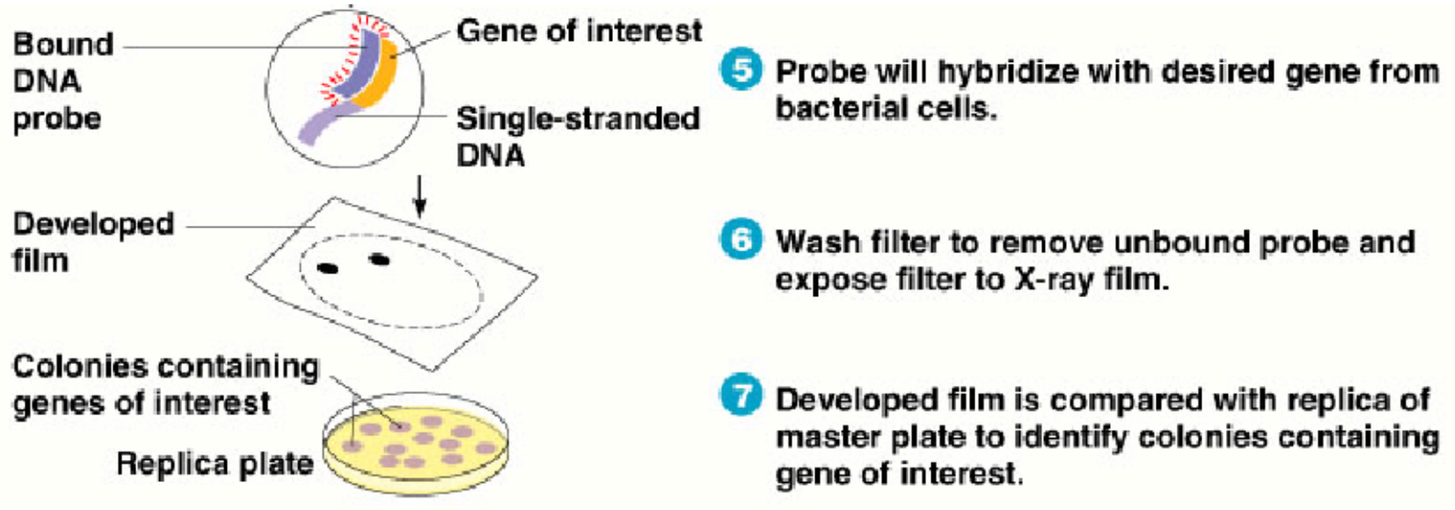
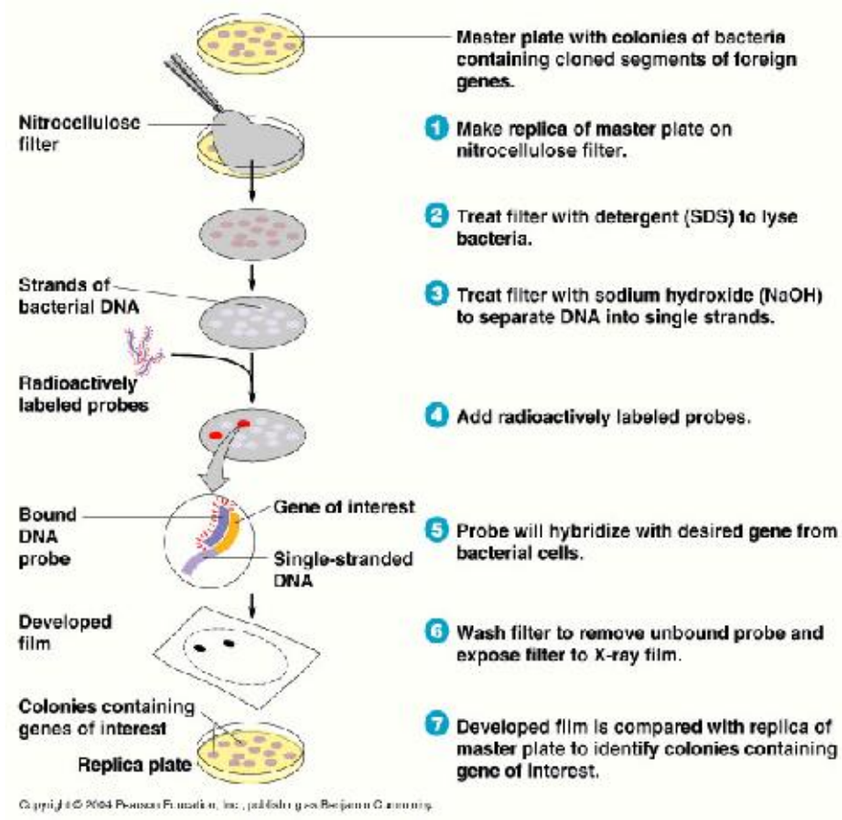


Figure 9.12.2

Colony hybridization works for finding a specific gene



E.coli

- Used because it is easily grown and its genomics are known
- Need to eliminate endotoxin from products
- Cells must be lysed to get product

DNA sequencing of a cloned piece of DNA

- Clone DNA to produce many copies to analyze sequence
- Can then be used analyze a person's DNA for the presence/absence of the gene
- Can be used to identify pathogenic strains of bacteria
- Shot gun sequencing
- Southern Blot

Random Shotgun Sequencing

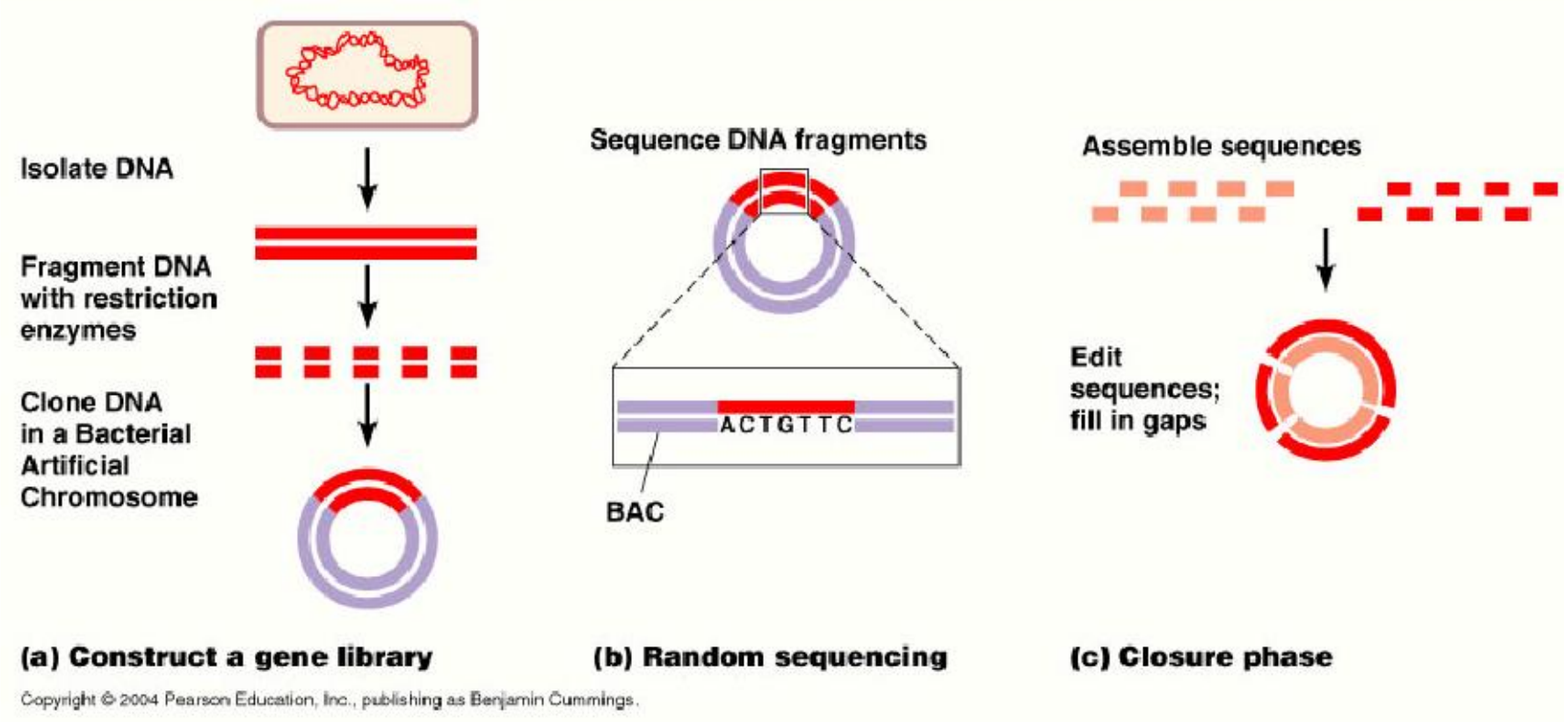
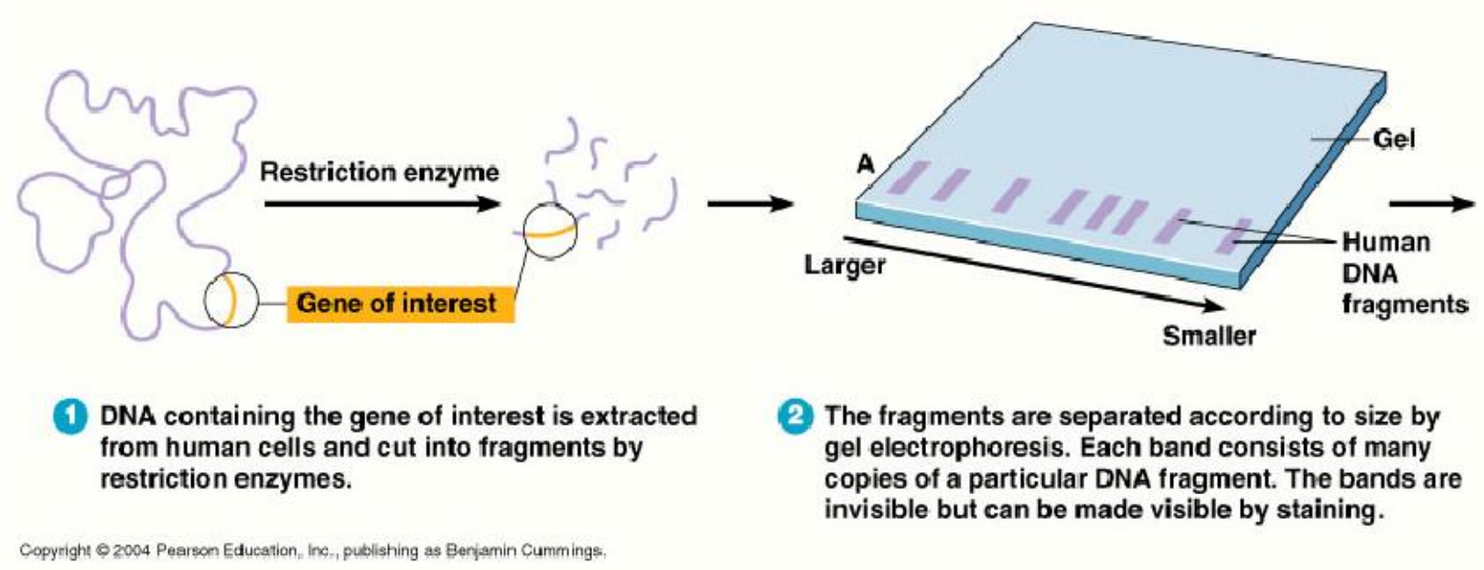


Figure 9.14

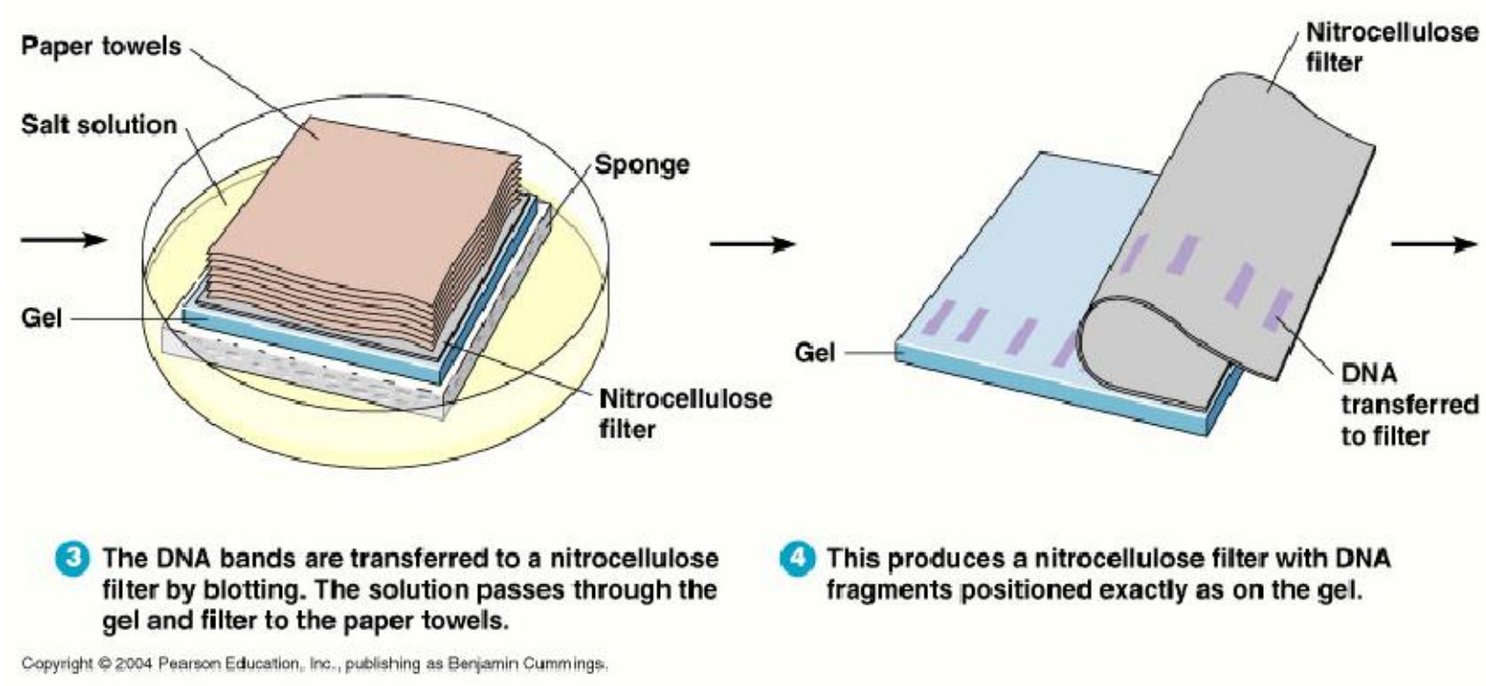
Southern Blot technique

- 1975 by Edward Southern
- Utilizes the idea of nucleic acid hybridization to target DNA
- DNA is cut into fragments with restriction enzymes
- Pieces of DNA are separated based on size on an agarose gel
- Probes are used to identify the target gene/sequence of DNA

Southern Blot technique



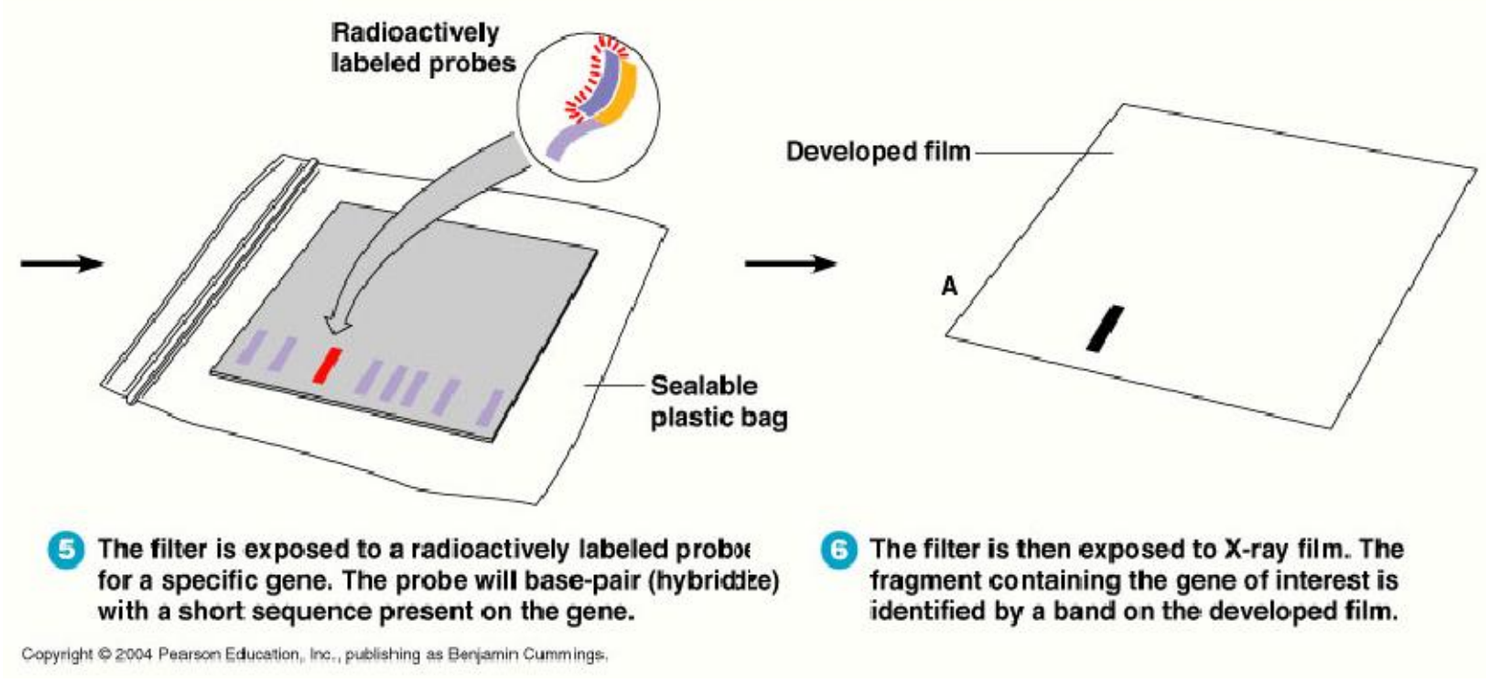
Southern Blot technique



3 The DNA bands are transferred to a nitrocellulose filter by blotting. The solution passes through the gel and filter to the paper towels.

4 This produces a nitrocellulose filter with DNA fragments positioned exactly as on the gel.

Southern Blot technique



Polymerase Chain Reaction

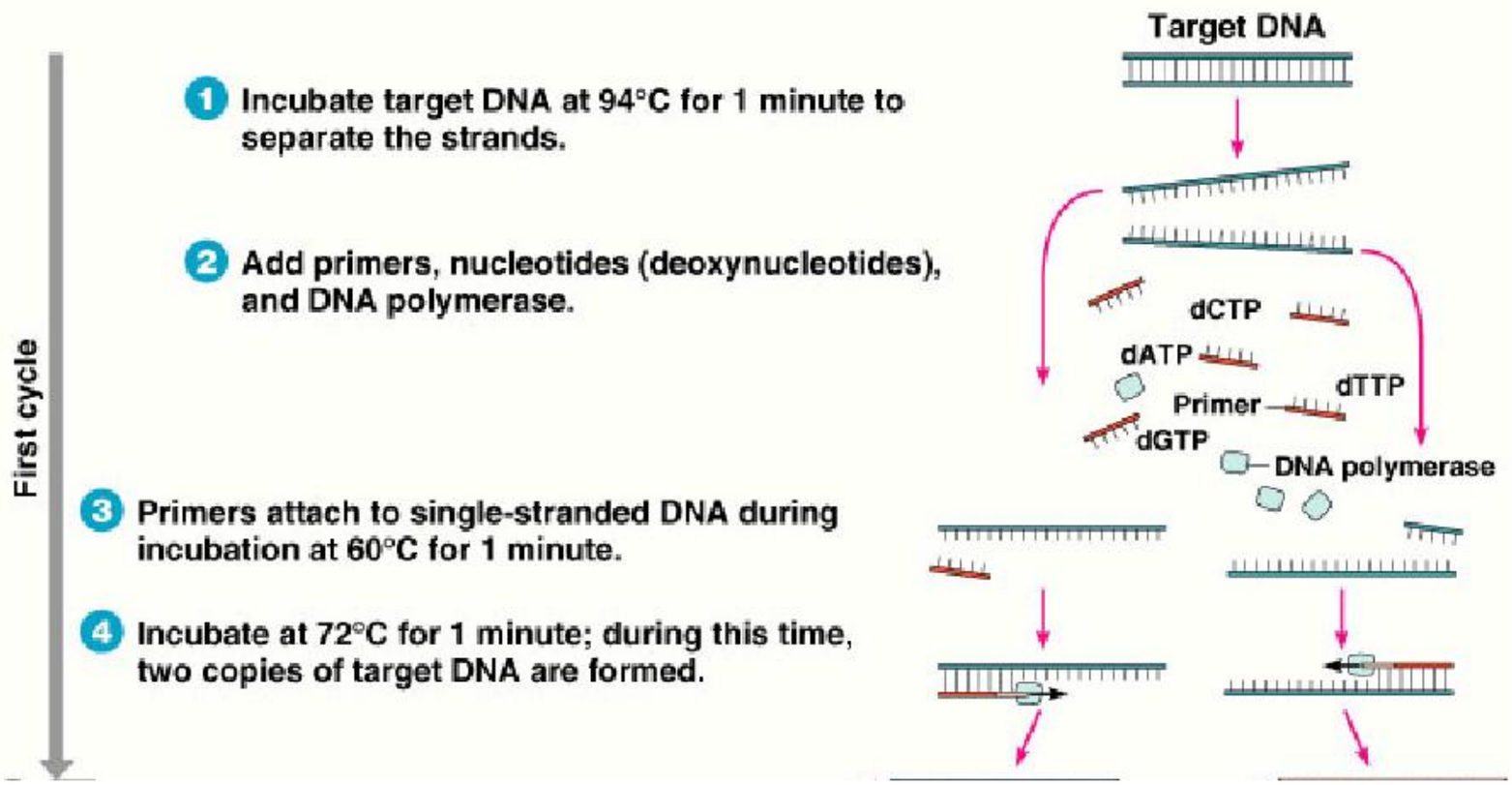


Figure 9.4.1

Polymerase Chain Reaction

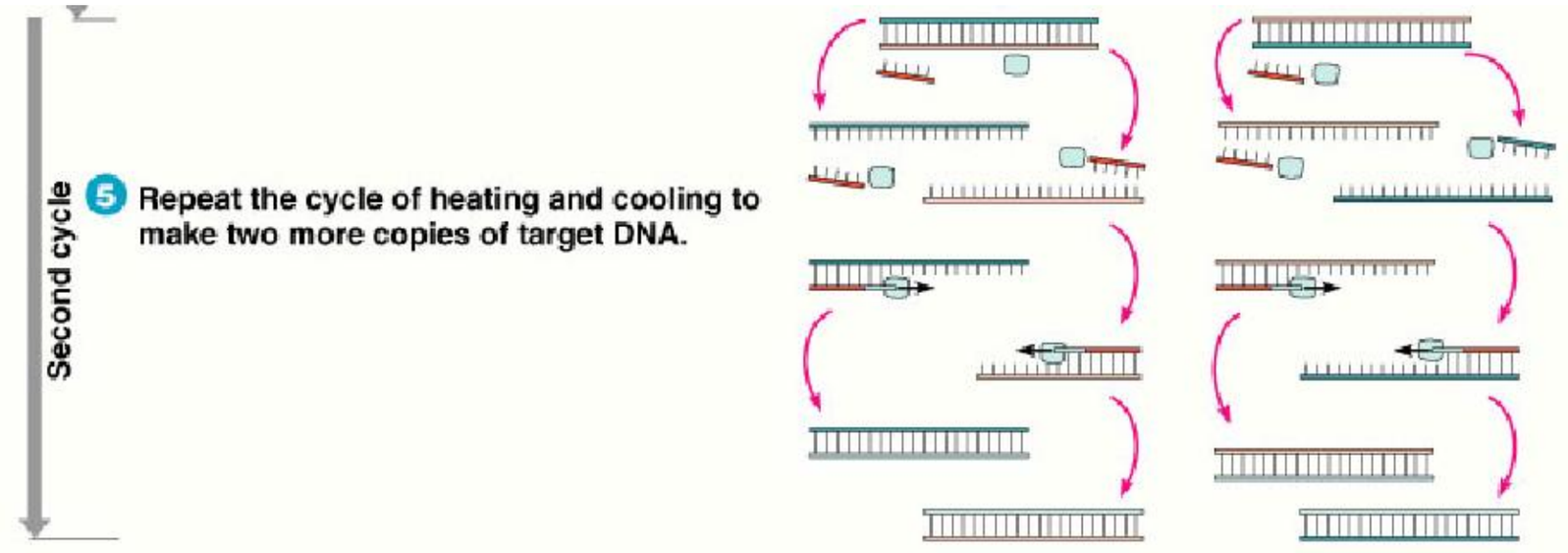


Figure 9.4.2

Polymerase Chain Reaction (PCR)

- A technique used to make more copies of DNA in vitro (enzymatically)
- Requires all the building blocks of DNA
- DNA Polymerase (Taq polymerase)
  - From thermophilic bacteria,

  Thermus aquaticus

  WHY?

Polymerase Chain Reaction (PCR)

- Used to
  - Clone DNA for recombination
  - Amplify DNA to detectable levels
  - Sequence DNA
  - Diagnose genetic disease
  - Detect pathogens

Therapeutic Applications

- Subunit vaccines
- Nonpathogenic viruses carrying genes for pathogen's antigens as vaccines
- Gene therapy to replace defective or missing genes
- Human Genome Project
  - Nucleotides have been sequenced
  - Human Proteome Project may provide diagnostics and treatments

Scientific Applications

- Understanding of DNA
- Sequencing organisms' genomes
- DNA fingerprinting for identification

DNA fingerprinting