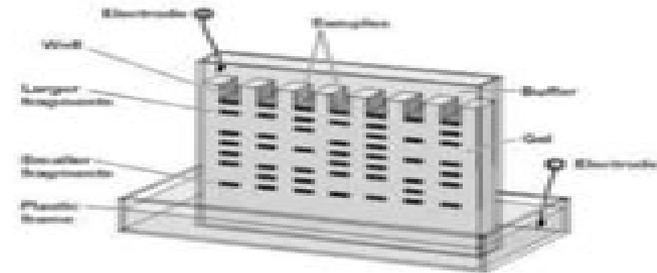# Manipulating DNA Notes

Scientists use their knowledge of the structure of DNA and its chemical properties to study and change DNA molecules.

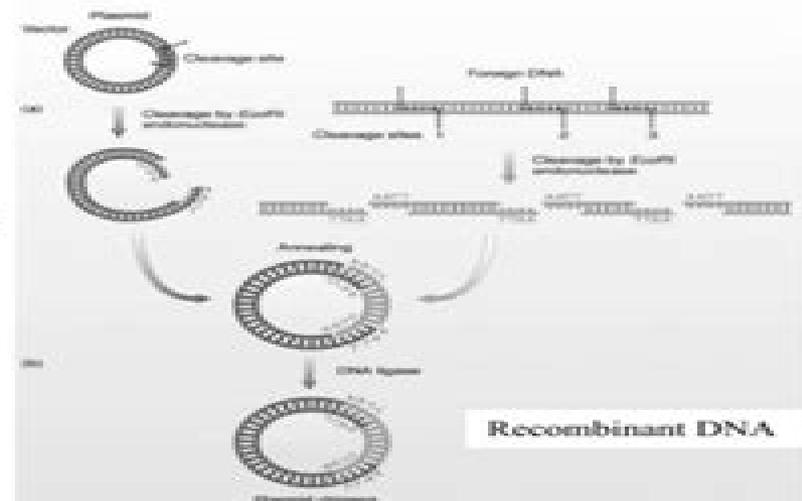**Tools of Molecular Biology**

1. Gel Electrophoresis
   a. DNA Extraction- using chemicals to remove DNA from the nucleus of the cell
   b. Cutting DNA- DNA is much too long to be analyzed, so we cut it into smaller fragments using restriction enzymes, which cut DNA at a specific sequence of nucleotides.
   c. Sequencing DNA- DNA fragments can be separated by weight and length in order to analyze specific sequences (order of bases).
      - DNA fragments are inserted into a porous gel, and then a voltage is passed through, pulling the DNA to the opposite side of the gel.
      - DNA is negatively charged.
      - The DNA sequence can then be "read" as the lighter segments will travel further and faster.

2. Transformation- a cell takes in DNA from another source and combines them together to make recombinant DNA.
   - Recombinant DNA- DNA that is made by connecting fragments of DNA from different sources.
   - Restriction enzymes are used to cut a sequence of DNA.
   - What is leftover? A double stranded DNA fragment with a single stranded end.
   - Next, a vector is used to insert the DNA into its new source. A vector is a carrier in which foreign DNA can be transferred into a host cell.
   - Vectors can be:
      a. Mechanical- to introduce DNA into a nucleus
      b. Biological- virus DNA or plasmids (a small ring of DNA found in a bacterial cell).
   - Finally, gene splicing rejoins the cut pieces of DNA through the use of DNA ligase (an enzyme which acts like tape to join the pieces together).

Recombinant DNA

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