

General introduction

LAB 1

Safety issues

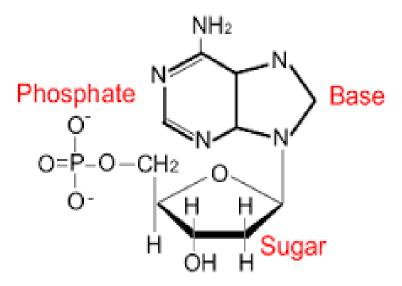
Major points:

- lab coat & gloves at all times, change when contaminated (for PCR fresh gloves/step)
- be familiar with eyewash station
- Dispose of everything as suggested
- Label clearly
- physical, biological & chemical hazards?

Nucleic acid

- macromolecule (monomeric nucleotides)
- carry genetic inf
- form structure
- all cells & viruses
- eg; DNA; RNA

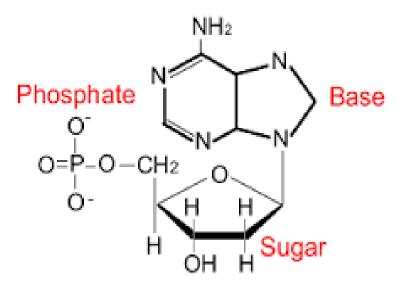
Nucleotide structure



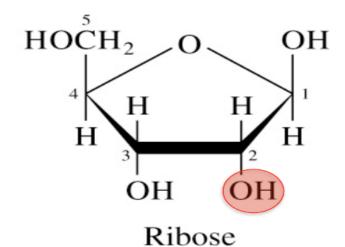
Nucleic acid (chemical structure)

- linear (eukaryotes), unbranched
- 3 components;
 - 1. phosphate backbone (PO4)⁻³
 - 2. pentose sugar
 - 3. nitrogenous base (purine or pyrimidine)

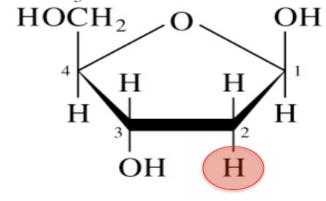
Nucleotide structure



Ribose and Deoxyribose sugars



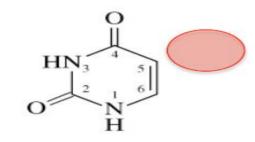
 $(\beta$ -D-Ribofuranose)



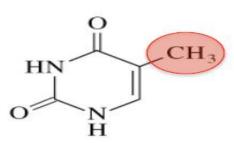
Deoxyribose (2-Deoxy-β-D-ribofuranose)

Nitrogenous base (nucleobases)

PYRIMIDINES



Uracil (2,4-Dioxopyrimidine)



Thymine (2,4-Dioxo-5-methylpyrimidine)

Cytosine (2-Oxo-4-aminopyrimidine)

PURINES

$$\begin{array}{c|c}
NH_2 \\
N & 5 \\
 & 7 \\
 & 8 \\
 & N
\end{array}$$

$$\begin{array}{c|c}
N \\
7 \\
8 \\
N \\
N \\
H
\end{array}$$

Adenine (6-Aminopurine)

$$H_{2} \stackrel{O}{\longrightarrow} N$$

$$N$$

$$N$$

$$N$$

$$N$$

$$N$$

$$N$$

$$N$$

Guanine (2-Amino-6-oxopurine)

Nucleic acid components

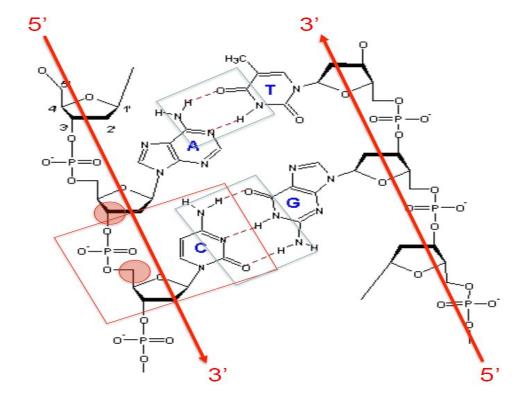
Nucleobases : C,G,T,A,U

Nucleosides (sugar + base)

Nucleotides (sugar + base + phosphate group)
 monomers of DNA, RNA

Base pairing

- 3'-5' phosphodiester link
- 2 strands liked by hydrogen bond
- stand run in opposite direction



Types of nucleic acids

DNA

Double stranded Single stranded

Ribose Deoxyribose (lack one O2 atom)

Thymine Uracil

Carry genetic information Protein synthesis

tRNA, mRNA, rRNA

DNA storage

Genes - DNA - chromatin - chromosomes

DNA make up of genes

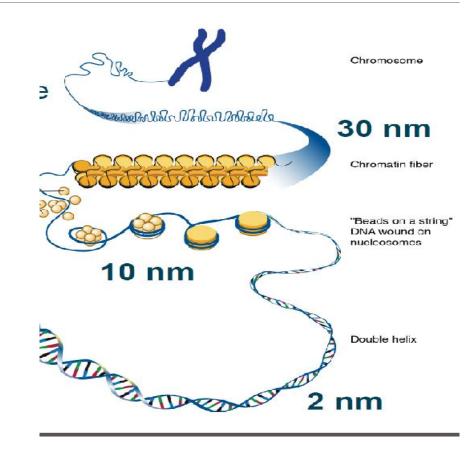


DNA + protein = chromatin

to maintain chromosome structure

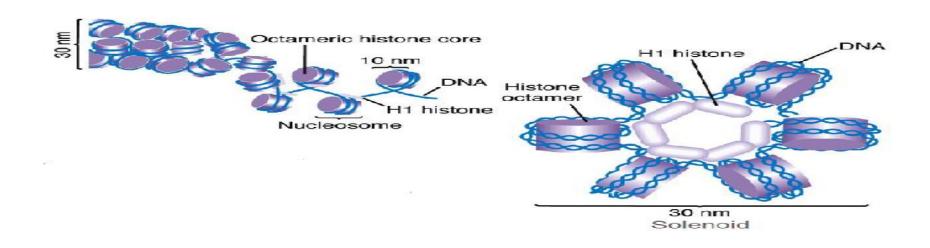


Chromosomes are made up of DNA



Histones

- +ve charged
- Compact & organize
- Control which part of DNA are transcribed



The human genome

- 3 billion bb.
- 200 volume the size of telephone
 Book to hold the information.

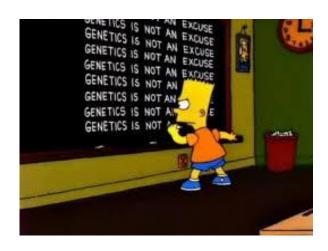
..... So compact structure



Genetics

study of the effect of genetic differences in organisms

study of the mutants organisms with respect to the wild-type (normal phenotype)



Molecular genetics

Study the structure and function of genes at a molecular level with understanding the interactions bet the various **systems** of a cell and learning how these interactions are regulated

Techniques:

Cloning

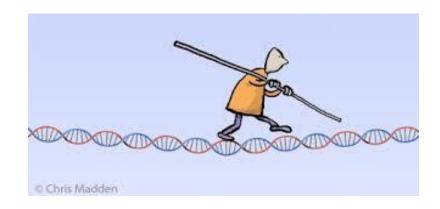
PCR

Gel electrophoresis

Sothern plot

Sequencing

Arrays



Cytogenetics

visual study of chromosomes at microscopic level

Techniques:

Karyotype (chromosomal complement)

Fish

Idiogram (stylised form of karyotype)



Mutation

- errors usually occur in the polymerization of the second strand
- error rate; 1 error / 10 100 million bases 'proofreadin function'
- mutagenic:

radiations, chemicals or inherited

THANKS FOR YOUR ATTENTION

ANY QUESTIONS?

