Introduction to genome biology

Sandrine Dudoit and Robert Gentleman

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Outline

- Cells and cell division
- DNA structure and replication
- Proteins
- Central dogma: transcription, translation
- Pathways









Cells

- Each cell contains a complete copy of an organism's **genome**, or blueprint for all cellular structures and activities.
- Cells are of many different types (e.g. blood, skin, nerve cells), but all can be traced back to a single cell, the fertilized egg.

Cell composition

- 90% water.
- Of the remaining molecules, dry weight
 - 50% protein
 - 15% carbohydrate
 - 15% nucleic acid
 - 10% lipid
 - 10% miscellaneous.
- By element: 60% H, 25% O, 12%C, 5%N.

The genome

- The genome is distributed along chromosomes, which are made of compressed and entwined DNA.
- A (protein-coding) gene is a segment of chromosomal DNA that directs the synthesis of a protein.

Eukaryotes vs. prokaryotes



Eukaryotes vs. prokaryotes

- Prokaryotic cells: lack a distinct, membrane bound nucleus.
 E.g. bacteria.
- Eukaryotic cells: distinct, membrane
- bound nucleus.
- Larger and more complex in structure than prokaryotic cells.
- E.g. mammals, yeast.



The eukaryotic cell

- Nucleus: membrane enclosed structure which contains chromosomes, i.e., DNA molecules carrying genes essential to cellular function.
- Cytoplasm: the material between the nuclear and cell membranes; includes fluid (cytosol), organelles, and various membranes.
- Ribosome: small particle composed of RNAs and proteins that functions in protein synthesis.

The eukaryotic cell

- **Organelle**: a membrane enclosed structure found in the cytoplasm.
- Vesicle: small cavity or sac, especially one filled with fluid.
- Mitochondrion: organelle found in most eukaryotic cells in which respiration and energy generation occurs.
- Mitochondrial DNA: codes for ribosomal RNAs and transfer RNAs used in the mitochondrion; contains only 13 recognizable genes that code for polypeptides.

The eukaryotic cell

- **Centrioles**: either of a pair of cylindrical bodies, composed of microtubules (spindles). Determine cell polarity, used during mitosis and meiosis.
- Endoplasmic reticulum: network of membranous vesicles to which ribosomes are often attached.
- Golgi apparatus: network of vesicles functioning in the manufacture of proteins.
- Cilia: very small hairlike projections found on certain types of cells. Can be used for movement.

The human genome

- The human genome is distributed along 23 pairs of chromosomes
 - -22 autosomal pairs;
 - -the sex chromosome pair, XX for females and XY for males.
- In each pair, one chromosome is paternally inherited, the other maternally inherited (cf. meiosis).





Chromosome banding patterns														
		2	3	4	5	6	7	8	9	10	11	12		
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DNA structure



"We wish to suggest a structure for the salt of deoxyribose nucleic acid (D.N.A.). This structure has novel features which are of considerable biological interest."

J.D. Watson & F. H. C. Crick. (1953). Molecular structure of Nucleic Acids. Nature. 171: 737-738.

DNA structure

- A deoxyribonucleic acid or DNA molecule is a double-stranded polymer composed of four basic molecular units called nucleotides.
- Each nucleotide comprises
 - a phosphate group;
 - a deoxyribose sugar;
 - one of four nitrogen bases:
 - purines: adenine (A) and guanine (G),
 - pyrimidines: cytosine (C) and thymine (T).

DNA structure

- Base pairing occurs according to the following rule:
 - C pairs with G,
 - A pairs with T.
- The two chains are held together by hydrogen bonds between nitrogen bases.

DNA structure











DNA strucure

- Polynucleotide chains are directional molecules, with slightly different structures marking the two ends of the chains, the so-called 3' end and 5' end.
- The 3' and 5' notation refers to the numbering of carbon atoms in the sugar ring.
- The 3' end carries a sugar group and the 5' end carries a phosphate group.
- The two complementary strands of DNA are antiparallel (i.e, 5' end to 3' end directions for each strand are opposite)

Genetic and physical maps

- Physical distance: number of base pairs (bp).
- Genetic distance: expected number of crossovers between two loci, per chromatid, per meiosis.
 Measured in Morgans (M) or centiMorgans (cM).
- 1cM ~ 1 million bp (1Mb).

The human genome in numbers

- 23 pairs of chromosomes;
- 2 meters of DNA;
- 3,000,000,000 bp;
- 35 M (males 27M, females 44M);
- 30,000 40,000 genes.

DNA replication



"It has not escaped our notice that the specific pairing we have postulated immediately suggests a possible copying mechanism for the genetic material."

J.D. Watson & F. H. C. Crick. (1953). Molecular structure of Nucleic Acids. Nature. 171: 737-738.





DNA replication

- In the replication of a double-stranded or duplex DNA molecule, both parental (i.e. original) DNA strands are copied.
- The parental DNA strand that is copied to form a new strand is called a **template**.
- When copying is finished, the two new duplexes each consist of one of the original strands plus its complementary copy **semiconservative** replication.





- Many enzymes are required to unwind the double helix and to synthesize a new strand of DNA.
- The unwound helix, with each strand being synthesized into a new double helix, is called the replication fork.
- DNA synthesis occurs in the 5' → 3' direction.















Enzymes in DNA replication

- 1. **Topoisomerase**: removes supercoils and initiates duplex unwinding.
- 2. Helicase: unwinds duplex.
- **3. DNA polymerase**: synthesizes the new DNA strand; also performs proofreading.
- 4. **Primase**: attaches small RNA primer to singlestranded DNA to act as a substitute 3'OH for DNA polymerase to begin synthesizing from.
- **5.** Ligase: catalyzes the formation of phosphodiester bonds.
- 6. Single-stranded binding proteins: maintain the stability of the replication fork.

DNA polymerase

- There are different types of polymerases, DNA polymerase III is used for synthesizing the new strand.
- DNA polymerase is a holoenzyme, i.e, an aggregate of several different protein subunits.
- DNA polymerase proceeds along the template and recruits free dNTPs (deoxynucleotide triphosphate) to hydrogen bond with their appropriate complementary dNTP on the template.
- The energy stored in the triphosphate is used to form the covalent bonds.
- DNA polymerase uses a short DNA fragment or primer with a 3'OH group onto which it can attach a dNTP.







- Proteins: large molecules composed of one or more chains of amino acids, polypeptides.
- Amino acids: class of 20 different organic compounds containing a basic amino group (-NH₂) and an acidic carboxyl group (-COOH).
- The order of the amino acids is determined by the **base sequence** of nucleotides in the **gene** coding for the protein.
- E.g. hormones, enzymes, antibodies.













Differential expression

- Each cell contains a complete copy of the organism's genome.
- Cells are of many different types and states E.g. blood, nerve, and skin cells, dividing cells, cancerous cells, etc.
- What makes the cells different?
- Differential gene expression, i.e., when, where, and how much each gene is expressed.
- On average, 40% of our genes are expressed at any given time.





The **expression** of the genetic information stored in the DNA molecule occurs in two stages:

- (i) transcription, during which DNA is transcribed into mRNA;
- (ii) **translation**, during which mRNA is translated to produce a protein.

DNA → mRNA → protein

Other important aspects of regulation: methylation, alternative splicing, etc.



RNA

- A ribonucleic acid or RNA molecule is a nucleic acid similar to DNA, but
 - single-stranded;
 - ribose sugar rather than deoxyribose sugar;
 - uracil (U) replaces thymine (T) as one of the bases.
- RNA plays an important role in protein synthesis and other chemical activities of the cell.
- Several classes of RNA molecules, including messenger RNA (mRNA), transfer RNA (tRNA), ribosomal RNA (rRNA), and other small RNAs.



- DNA: sequence of four different nucleotides.
- Proteins: sequence of twenty different amino acids.
- The correspondence between DNA's fourletter alphabet and a protein's twenty-letter alphabet is specified by the genetic code, which relates nucleotide triplets or codons to amino acids.



The Genetic Code

Mapping between codons and amino acids is many-to-one: 64 codons but only 20 a.a..

Third base in codon is often redundant, e.g., stop codons.



Transcription

- · Analogous to DNA replication: several steps and many enzymes.
- RNA polymerase synthesizes an RNA strand complementary to one of the two DNA strands.
- The RNA polymerase recruits rNTPs (ribonucleotide triphosphate) in the same way that DNA polymerase recruits dNTPs (deoxunucleotide triphospate).
- However, synthesis is single stranded and only proceeds in the 5' to 3' direction of mRNA (no Okazaki fragments).

Transcription

- The strand being transcribed is called the template or antisense strand; it contains anticodons.
- The other strand is called the sense or coding strand; it contains codons.
- The RNA strand newly synthesized from and complementary to the template contains the same information as the coding strand.

Transcription

- **Promoter.** Unidirectional sequence upstream of the coding region (i.e., at 5' end on sense strand) that tells the RNA polymerase both **where** to start and on **which strand** to continue synthesis. E.g. TATA box.
- Terminator. Regulatory DNA region signaling end of transcription, at 3' end .
- **Transcription factor.** A protein needed to initiate the transcription of a gene, binds either to specific DNA sequences (e.g. promoters) or to other transcription factors.



Exons and introns

- Genes comprise only about 2% of the human genome.
- The rest consists of non-coding regions
 - chromosomal structural integrity,
 - cell division (e.g. centromere)
 - regulatory regions: regulating when, where, and in what quantity proteins are made .
- The terms exon and intron refer to coding (translated into a protein) and non-coding DNA, respectively.

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Alternative splicing

- There are more than 1,000,000 different human antibodies. How is this possible with only ~30,000 genes?
- Alternative splicing refers to the different ways of combining a gene's exons. This can produce different forms of a protein for the same gene.
- Alternative pre-mRNA splicing is an important mechanism for regulating gene expression in higher eukaryotes.
- E.g. in humans, it is estimated that approximately 30% of the genes are subject to alternative splicing.



Immunoglobulin

- B cells produce antibody molecules called immunoglobulins (Ig) which fall in five broad classes.
- Diversity of Ig molecules – DNA sequence: recombination,
 - DNA sequence: recombination mutation.
 - mRNA sequence: alternative splicing.
 - Protein structure: post-translational proteolysis, glycosylation.



lgG1

Post-translational processing

- Folding.
- Cleavage by a proteolytic (protein-cutting) enzyme.
- Alteration of amino acid residues
 - phosphorylation, e.g. of a tyrosine residue.
 - glycosylation, carbohydrates covalently attached to asparagine residue.
 - methylation, e.g. of arginine.
- Lipid conjugation.

Functional genomics

 The various genome projects have yielded the complete DNA sequences of many organisms.

> E.g. human, mouse, yeast, fruitfly, etc. Human: 3 billion base-pairs, 30-40 thousand genes.

• Challenge: **go from sequence to function**, i.e., define the role of each gene and understand how the genome functions as a whole.

Pathways

- The complete genome sequence doesn't tell us much about how the organism functions as a biological system.
- We need to study how different gene products interact to produce various components.
- Most important activities are not the result of a single molecule but depend on the **coordinated effects** of multiple molecules.

TFG- β pathway

- Transforming Growth Factor beta, TGF-β, plays an essential role in the control of development and morphogenesis in multicellular organisms.
- The basic pathway provides a simple route for signals to pass from the extracellular environment to the nucleus, involving only four types of molecules.





TFG- β pathway

- Extracellular TGF-β ligands transmit their signals to the cell's interior by binding to type II receptors, which form heterodimers with type I receptors.
- The receptors in turn activate the SMAD transcription factors.

TFG- β pathway

- Phosphorylated and receptor activated SMADs (R SWADs) form heterodimers with common SMADs (co SWADs) and translocate to the nucleus.
- In the nucleus, SMADs activate or inhibit the transcription of target genes, in collaboration with other factors.

Pathways

- http://www.grt.kyushu u.ac.jp/spad/
- There are many open questions regarding the relationship between gene expression levels (e.g. mRNA levels) and pathways.
- It is not clear to what extent microarray gene expression data will be informative.

WWW resources

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