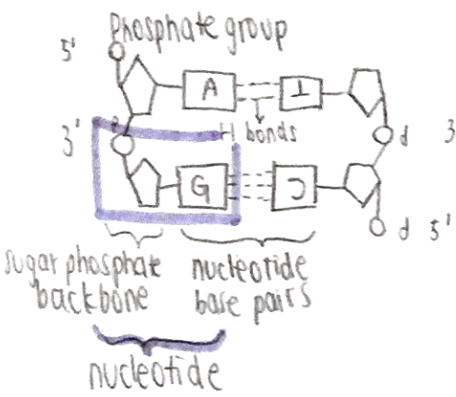


Molecular Genetics

Basics: Structure



Nitrogenous bases

Purine: Adenine, Guanine

Pyrimidine: Thymine, Cytosine

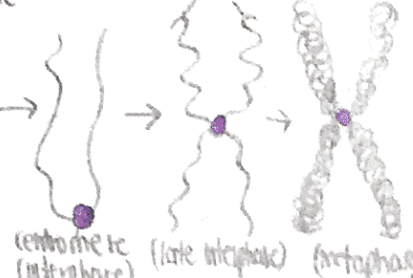
go through complementary base pairing

segment of DNA containing genetic info to make a protein

sugar phosphate backbone



2 copies of DNA



Gene $\xrightarrow{\times n}$ DNA $\xrightarrow{+ \text{histones}}$ Chromatin \rightarrow Chromosome

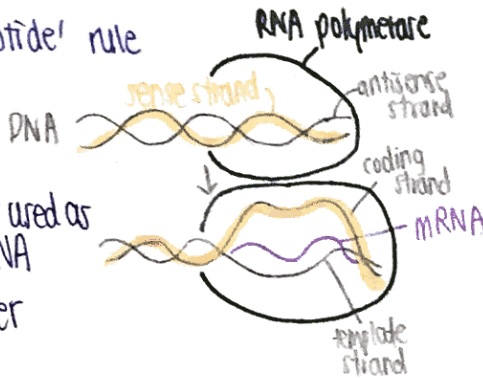
Tri-nucleotide sequences (codon) \rightarrow A-A- for both DNA & mRNA

Protein synthesis

'One gene one polypeptide' rule

TRANSCRIPTION

- DNA section unwinds
- one DNA strand (out of 2) is used as template to transcribe mRNA
- mRNA small enough to enter cytoplasm



messenger RNA
Transfer RNA
Ribosomal RNA
Ribonucleic acid

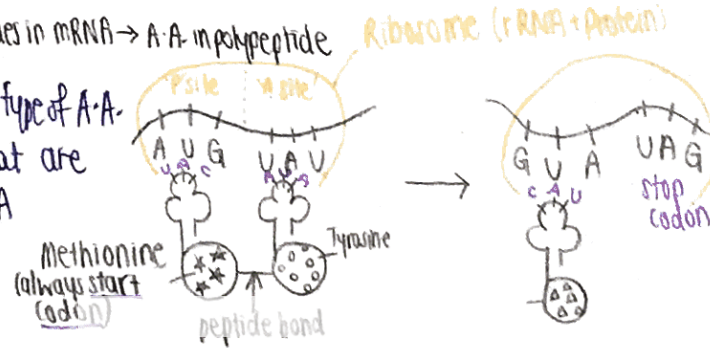
DNA helicase unwinds DNA by breaking H-bonds (own bases)

VS (RNA polymerase reads template strand & transcribes on pre-mRNA molecule) (RNA)

TRANSLATION

conversion of nucleotides in mRNA \rightarrow A-A in polypeptide

- Each type of tRNA is specific \rightarrow one type of A-A.
- tRNA has 3 nucleotides (anti-codon) that are complementary to codon on mRNA

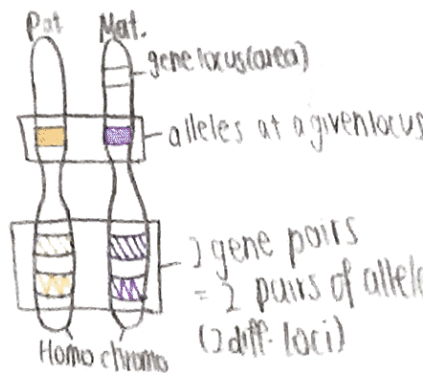


Ribosome (rRNA + Protein)

- ① mRNA codons complementary to tRNA anti-codons
- ② Peptide bonds bwn 2 A-A.
- ③ Ribosome moves down 3 bases/nucleotides to form more A-A & peptide bonds
- ④ Upon reaching stop codon: UAA, UAG, UGA, ribosome detaches from mRNA to go to other mRNA or G.A. to be repackaged etc.

Cell division

- Homologous chromosome \rightarrow A pair of chromo. of same length & sequence of genes (homo chromo) \rightarrow Paternal & maternal
- 1st 22 pairs chromo \rightarrow Autosomal, last \rightarrow sex chromo arranged by decreasing length \uparrow
- Alleles are alternative forms of a gene (of same relative posⁿ) on a pair of homologous chromosomes

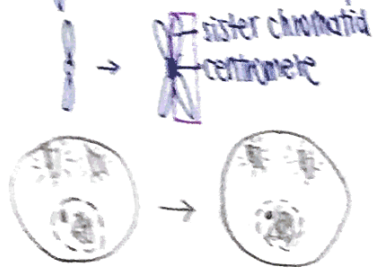


Mitosis

2 daughter nuclei each contains same no. of chromo as parent nucleus

Interphase

- Chromatins present, distinguished indiv.
- DNA replicat^d
- Centrioles in animals, centrosomes (MTOC - microtubule organization center) in plants

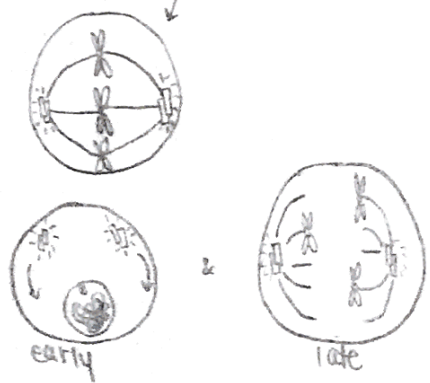


Prophase (break-down & form)

- duplicated chromatins condense & coil → chromo.
- Early prophase → distinct nuclear envelope
- Asters form and centrioles (microtubules) move to opp. poles of cell
- Nuclear envelope disappears
- Late prophase → Nuclear envelope disintegrates
- Spindle fibers form & extend

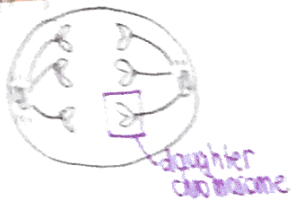
Metaphase (line up)

- Chromo. line up along equator
- Held in place by spindle fibers at centromere



Anaphase (separate)

- Centromeres separate as spindle fibers shorten & pull sister chromo. to opp. poles



Telophase (opp. of Prop)

- Daughter chromo reach poles
- Spindle fibers break down & nuclear envelope forms ad. chromo. at ea. pole
- Nuclear forms & chromo. → chromatin



Cytokinesis

- Division of cytoplasm
- Cleavage furrow appears in cytoplasm b/w 2 nuclei
- deepens & forms 2 daughter cells

plants

- Vesicles from Golgi A. line up at equator of spindle
- Fuse to form cell plate
- cell plate elongates & fuse w/ cell wall
- partitions cell into 2



Meiosis I

4 daughter nuclei each contains half the no. of chromo as parent nucleus

Interphase

same as mito! → Duplicate

Prophase I

- Chromatin → Chromo.
- Homologous chromo. pair up along their length
- Centrioles move to opp. poles
- Crossing over may take place: two sister chromo. break and exchange chromatid seg. info

Metaphase I

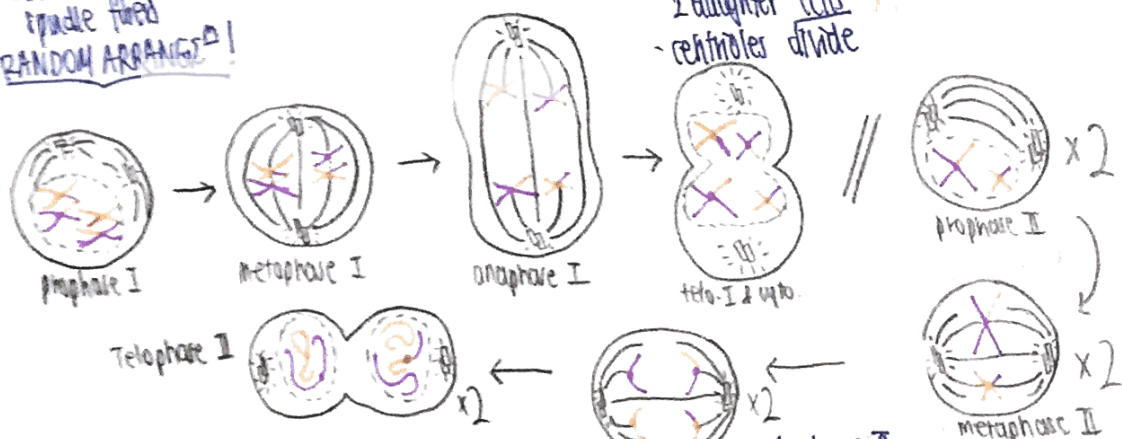
- Pairs of homo chromo line up along equator
- centromere attached to spindle fibers
- RANDOM ARRANGEMENT!

Anaphase I

- Homo chromo pair separate, pulled to opp. poles
- no splitting of centromeres!

Telophase I

- spindle fibers disintegrate
- Nuclear envelope forms and chromo. divide
- 2 daughter cells



Prophase II

no DNA replicat^d → 4 daughter cells formed (n)

Metaphase II

Anaphase II

Telophase II

Mitosis

Importance

- Growth of organism
- Repair of worn out parts of body
- Healing wounds
- Asexual reproduction

Meiosis

- Produce haploid gametes
 - ↳ maintains diploid no. in species
- Genetic variation
 - ↳ Chromatids formed during Pro. I (new combi. of alleles)
 - ↳ Ind. assortⁿ & segregatⁿ of homo. chromo during Meta. I (4 diff. gametes prod. & 4 possible combi.)
- Random fusion of gametes
 - ↳ chances of survival, favourable characteristics, changing env.

Pairing up of homo chromo

X

✓

Crossing over & chiasmata

X

✓

Arr. of chromo.

Single row

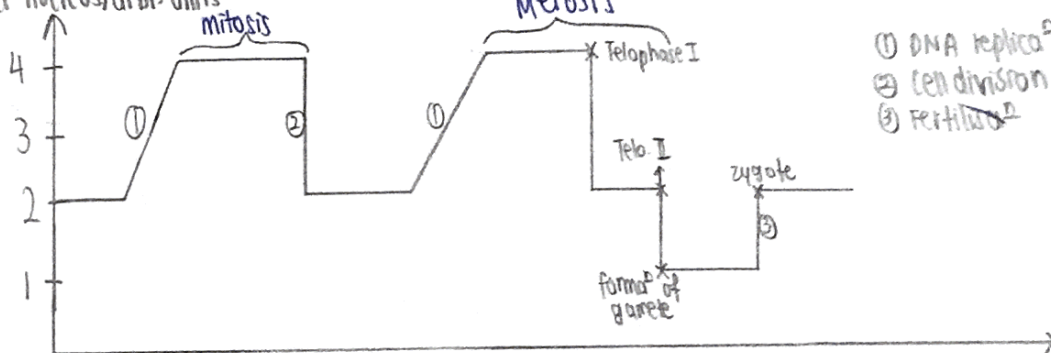
2 rows (Meta. I)

Separatⁿ of chromo

Chromatids move to opp. poles during Ana

Chromo. move during Ana I
Chromatids Ana II

DNA per nucleus/cell. units



INHERITANCE

Genetic Cross diagram

Parental phenotype: Tall x Tall
genotype: Tt x Tt
gametes: T, t, T, t
offspring's genotype: TT, Tt, Tt, tt
phenotype: Tall, Tall, Tall, Short
ratio: 3:1

Punnett's Sq

	T	t
T	TT	Tt
t	Tt	tt

Examples of terms

Homozygous	AA/aa
Heterozygous	Aa
	dominant (A) / recessive (a)
Chance	75% or $\frac{3}{4}$
Probability	$\frac{3}{4}$

* Expected ratios from genetic crosses will not occur as exp. in small sample sizes.

Ratio based on chance & probability only $n > 100$, then ratio appears

* Co-dominance: Both alleles fully expressed (eg. Purple & white flower)
Incomplete dominance: Blend of alleles (eg. light purple flower)

Blood Types

A: $I^A I^A$, $I^A i$
B: $I^B I^B$, $I^B i$
AB: $I^A I^B$
O: $i i$

* A & B are mutaⁿ of O!

Mutation (spontaneous change in gene structure, chromo or chromo no.)

- May be inheritable, lethal or beneficial
- Somatic mutaⁿ (body cells) may be inheritable/cancer
- mutaⁿ during gamete prodⁿ inherited by offspring
- Produces genetic variaⁿ

CASE STUDY: Sickle cell anaemia

- HbS instead of HbA (autosomal recessive)
- HbS mol clump pt. → sickle shape, reduce O_2 transporting efficiency, RBC more fragile & block blood vessels. survives only 10-20 days, ↓ blood count, → anaemia
- GENETIC MUTAⁿ

#2 Case Study: Down's syndrome

- Change in chromo. no., extra copy of chromo. 21
- Older mom, ↑ risk
- Happens during gamete prod
→ nondisjunction (Anaphase I/II)
chromo. → Chromatids

Factors causing mutation

- mutagens are UV light, radiaⁿ & chem.

q. misheard par
turns blonde
LSD

Autosomal recessive (non-sex chromo must both appear dominant only 1)