

Inheritance

Charles Darwin



Inheritance

Inheritance is the process by which genetic information is passed on from parent to child **Hereditary traits** can be passed down to you by your parents:



ability to roll tongue



allergy



ability to taste PTC (bitter compound)



dimples



right or left handed



attached or free earlobe



(a) Distinguish between the terms gene and allele

(b) Explain the terms dominant, recessive, codominant, homozygous, heterozygous, phenotype and genotype



Wast-

Gene	A sequence of DNA nucleotides that stores information used to
	make a polypeptide
	A unit of inheritance passed from parents to offspring
	Each gene occupies a locus, a specific position on chromosome
Allele	• Alleles are different forms of a gene that leads to variations in those traits
	Alleles arises due to mutation
	• Alleles of a gene occupy the same locus on a pair of homologous
	chromosomes.
	• Individuals inherit two alleles for each gene, one from each parent.
	• For example, for the gene controlling eye color, there might be alleles for
	brown eyes and alleles for blue eyes.



Chromosome	 Chromosome is a compact structure visible in the nucleus during cell division and it is made up of DNA The place on the chromosome where the gene is located is called the gene locus.
Dominant	• A dominant allele is the allele that is always expressed in the phenotype, no matter under homozygous or heterozygous condition.
Recessive	A recessive allele is the allele that is only expressed under homozygous recessive condition.
Codominant	 When both alleles have an equal effect on the phenotype of the offspring. Both alleles are expressed in the phenotype.
Homozygous	• When organisms have two identical alleles of a particular gene. Allele can be either both dominant or both recessive.
Heterozygous	When organisms have two different alleles of a particular gene.
Genotype	 The genetic makeup of an organism. An organism's genotype is homozygous for a trait if the two alleles controlling the trait are identical, heterozygous for a trait if the alleles controlling the trait are different.
Phenotype	Phenotype refers to the observable, physical traits that are determined by the genotype.



(c) Predict the results of simple crosses with expected ratios of 3:1 and 1:1, using the terms homozygous, heterozygous, F1 generation and F2 generation

(d) Use genetic diagrams to solve problems involving monohybrid inheritance



Monohybrid Inheritance

Parental Generation



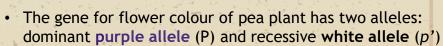


F1 Generation



F2 Generation





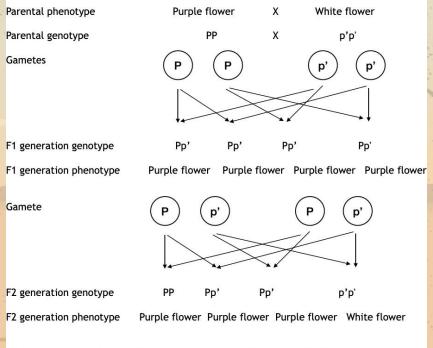
• When a homozygous purple plants (PP) is crossed with a homozygous white plant (p'p'), each organism inherits one allele from the mother and one allele from the father during sexual reproduction

- The offspring generation consisted of all purple-flowered plants even though their genotype is heterozygous (Pp') as the dominant purple allele (P) is expressed over recessive white allele (p')
- Self-pollination in the F1 generation produced a F2 generation where the phenotypic ratio of purple-flowered to white-flowered plants is 3:1



Monohybrid Inheritance

Let P represent the allele for purple flowers Let p' represent the allele for white flowers



- The gene for flower colour of pea plant has two alleles: dominant purple allele (P) and recessive white allele (p')
- When a homozygous purple plants (PP) is crossed with a homozygous white plant (p'p'), each organism inherits one allele from the mother and one allele from the father during sexual reproduction
- The offspring generation consisted of all purple-flowered plants even though their genotype is heterozygous (Pp') as the dominant purple allele (P) is expressed over recessive white allele (p')
- Self-pollination in the F1 generation produced a F2 generation where the phenotypic ratio of purple-flowered to white-flowered plants is 3:1



(e) Explain why observed ratios often differ from expected ratios, especially when there are small numbers of progeny





3 purple flower: 1 white flower is the expected ratios, but the actual observed ratio can be different especially when there are small numbers of progeny, because

- Fertilisation of the ova and sperms is a random event.
- Therefore the expected ratios are only based on chance and probabilities.

With small number of offspring, the observed ratios often differ from expected ratios. But, with large number of offspring (large sample size), the observed ratios will be closer to expected ratios.

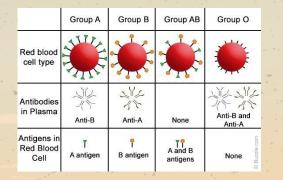


(f) explain codominance and multiple alleles with reference to the inheritance of the ABO blood group phenotypes (A, B, AB, O) and the gene alleles (I^A , I^B and I^O)





Codominance ABO blood group



Blood group		Homozygous	Heterozygous		
	phenotype	Genotype	Genotype		
	А	I ^A I ^A	IAIO		
	В	I ^B I ^B	I _B I _O		
AB		I ^A I ^B			
	0	I ^C	Io		

Complete dominance is when the heterozygote has the same phenotype as the dominant homozygote. In pea plant flower example, the recessive allele present in the heterozygote is masked by the dominant allele. This is because when you cross PP with pp, all the offspring in the F1 generation will have a genotype of Pp, and would therefore display purple flower phenotype.

Co-dominance is when both alleles contribute equally to the phenotype.

ABO blood group is determined by 3 alleles:

- IA: Allele for the production of Type A antigen (Blood Group A)
- I^B: Allele for the production of type B antigen (Blood Group B)
- I⁰: Allele that produces neither antigen (Blood Group O)

I^A and I^B are codominant, while I^O is recessive to both
For I^AI^B genotype, both antigen A and antigen B are expressed since they are codominant and each of the alleles produces its own antigen. Both alleles contribute to the phenotype, which is blood group AB.

• For I^AI^O and I^BI^O genotype, I^O is recessive to I^A and I^B thus, the phenotype is blood group A and B respectively.



Codominance ABO blood group example

			and set of the set of	
Parental phenotype	blood g	group B	X blood g	roup B
 Parental genotype		вю	X I ^B	ю
Gametes	ГВ	lo	ГВ	lo
			\ll	
	••			2004 - 20 4
F1 generation genotype	 В В	вю	Івіо	lolo
F1 generation phenotype	blood group B	blood group B	blood group B	blood group O

Parents with blood group B can produce offsprings with blood group O

F1 generation phenotypic ratio

3 blood group B : 1 blood group O



(g) describe the determination of sex in humans - XX and XY chromosomes



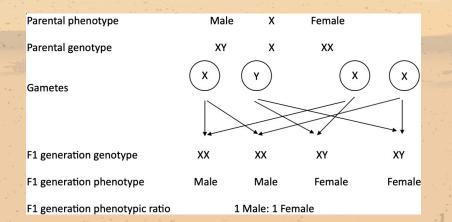


Sex determination

11 12 11 71 17 13 21 19

- A human cell has 23 pairs of chromosomes and the last pair is the sex chromosomes
- In humans, sex is determined by sex chromosomes.
- Human sex chromosomes are the X chromosome and the Y chromosome.
- X chromosome is much larger than the Y chromosome.
- Human males have one X chromosome and one Y chromosome (XY genotype)
- Human females have two X chromosomes (XX genotype)

Example of sex determination cross: equal probability of a male or female offspring





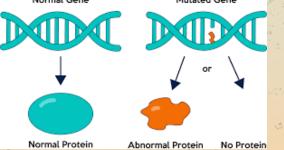
(h) describe mutation as a change in the sequence of a gene such as in sickle cell anaemia, or in the chromosome number, such as the 47 chromosomes in the condition known as Down syndrome

(i) name ionising radiation (e.g. X-ray) and chemical mutagens as factors which may increase the rate of mutation





Normal Gene Mutated Gene



1. Mutation is a random change in the :

- 1) structure/ sequence of a gene (gene mutation)
- 2) chromosome number

1. Mutations that take place in body cells other than gametes are called somatic mutations, which will not be passed on to the next generation

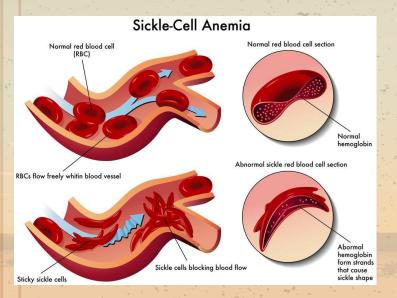
1. Mutation is spontaneous and can occur during replication of DNA

1. Mutagen increase the rate of mutation

1. Examples of mutagens:

- Ionising radiation: Ultraviolet radiation, x rays, gamma rays
- Chemical mutagens: such as benzene, ethidium bromide

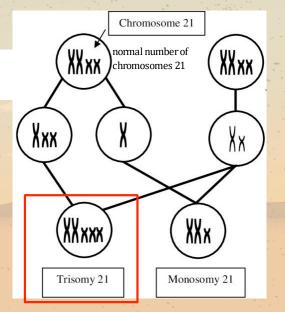
Gene mutation



- Sickle-cell anaemia is caused by a change in the sequence of nucleotides coding for the protein, haemoglobin
- It is a recessive condition, which means mutated allele only express in a homozygous recessive condition
- Heterozygous individual with one normal allele, one mutated allele are healthy but are carriers
- Normal red blood cells are flexible and can change their shape in order to pass through capillaries.
- Mutated gene produces Haemoglobin S (HbS) that tend to clump together, which result in sickle-shaped red blood cells that can block capillaries
- When oxygen concentration in the blood drops, the red blood cells become sickle-shaped and this lowers their surface area to volume ratio for diffusion of oxygen.
- Hence, they cannot transport oxygen as effectively as the normal red blood cells.
- In summary: changes in the sequence of nucleotides impacts the structure of proteins and consequently, the function of the protein



Chromosomal mutation



down syndrome is also known as trisomy 21

- Down syndrome is a condition caused by a chromosome mutation during meiosis (gamete production)
- The gamete has 2 copies of chromosome 21, thus upon fertilisation, the zygote inherits 3 copies of chromosome 21 and a total of 47 chromosomes
- This mutation is present in all body cells due to mitosis during zygote development.
- This chromosome mutation is far more likely to occur during ovum production than during sperm production.
- Women above 30 have a higher risk of carrying babies with Down syndrome.



(j) distinguish between continuous and discontinuous variation and give examples of each

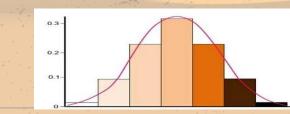






Variations are differences in traits between individuals of the same species.

	Discontinuous variation	Continuous variation	
Phenotype	Few clear-cut phenotypes with no intermediate	Range of phenotypes.	
Environment	Rarely affected by	Greatly affected by environmental	
influence	environmental conditions	conditions.	
Genes	Controlled by one or few genes	Controlled by many genes	
Additive effect	Not present	The effect of many genes add together and contribute the	
		phenotype	
Graph	Discrete groups	Normal distribution	



Height, skin colour, weight, intelligence

Examples

Blood groups, eyelid, flower colour in pea plant

Α

Number of people

Four distinct blood groups

B AB Blood group 0



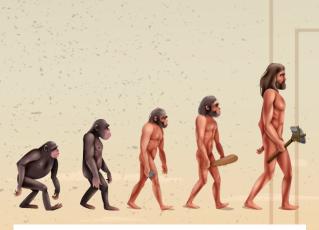
(k) State that variation and competition lead to differential survival of, and reproduction by, those organisms best fitted to the environment

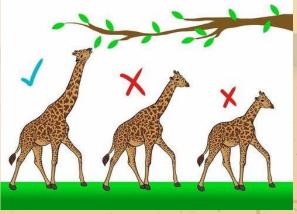
(l) give examples of environmental factors that act as forces of natural selection

(m) explain the role of natural selection as a possible mechanism for evolution which is a gradual change in the inheritable characteristics of a population over time

Natural selection

- . There are variation among individuals within the population such as giraffe with short and long neck. This is due to **environmental factors that act as forces of natural selection:** variation in climate, predation pressure, food availability, diseases and pathogens, geographic features, competition for resources
- 2. This therefore leads to **natural selection**, where individuals traits that are favoured by the environment / adapted to their environment are more likely to survive and reproduce and pass down their favourable alleles to their offspring
- 3. Overtime, offspring with favourable alleles and traits would increase in proportion
- 4. This leads to **evolution:** the gradual change in allele frequency in a population over generations. This could give rise to a new species.





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LEADERS IN THE CHANGING EDUCATION LANDSCAPE

FEATURED ON STRAITS TIMES

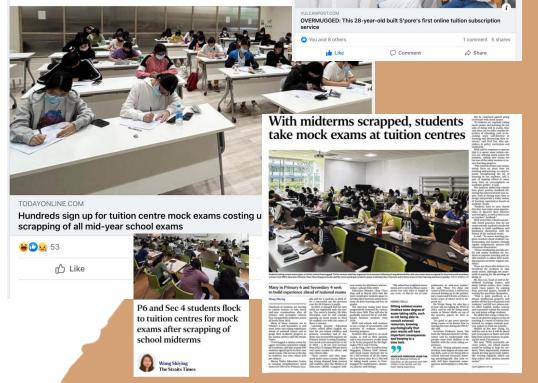
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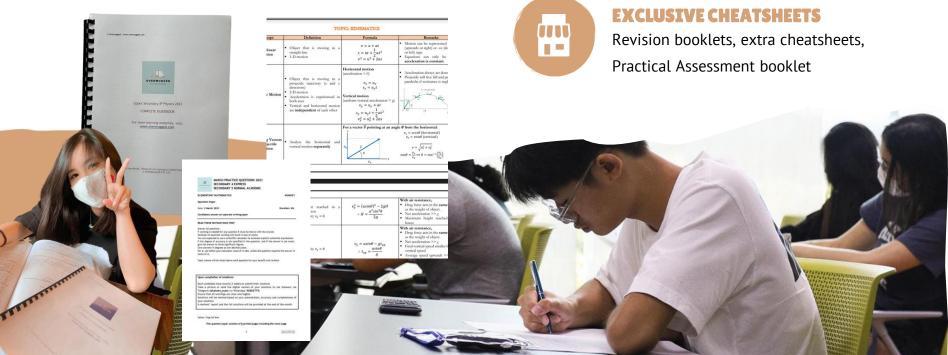
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TERM 1: NOV – JAN

Topical Recaps

Key highlight: Christmas Party

TERM 2: FEB – APR

Topical Mastery

Key highlight: March Holiday Cohesion Program

TERM 3: MAY – JUL

Prelim/EOY Preparation

Key highlight: Mock Prelim/EOY

TERM 4: AUG – OCT

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Key highlight: Mock Exams, Science Practical Assessment



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