

Definitions and Concepts for WJEC (Wales) Biology GCSE

Topic 2.3: DNA and Inheritance

Definitions in **bold** are for higher tier only

Definitions marked by "are for separate sciences only

Allele - A version of a gene (also known as variant).

Amino acids - Small molecules that make up a protein.

Chromosome - A long, coiled molecule of DNA that carries genetic information in the form of genes.

Complementary base pairing - Describes how the bases of DNA pair up with each other. A pairs with T; C pairs with G.

Cystic fibrosis - A genetic disorder caused by homozygous recessive alleles which causes mucus build up in the lungs.

DNA (Deoxyribonucleic acid) - 2 chains consisting of alternating sugar and phosphates which are twisted together and connected by bases (A, T, C, and G, **or adenine, thymine, cytosine and guanine**). The order of the bases determines the order of amino acids in a protein synthesised from that gene; the order of the amino acids linked together determines the structure of the synthesised protein. **+**

Disease - An illness that affects animal or plant health.

Dominant - An allele that is always expressed when present. It is represented by a capital letter.

Enzymes - Biological catalysts that increase the rate of chemical reactions.

F1 generation - The first generation of a genetic cross as a result of 2 organisms interbreeding.

F2 generation - The second generation of a genetic cross as a result of 2 organisms from the F1 generation interbreeding.

Fertilisation - When the male and female gametes fuse restoring the full chromosome number.

Gamete - Sex cells (sperm and egg cells) with half the usual number of chromosomes. They are involved in reproduction.

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Gene - A section of DNA that codes for a specific amino acid sequence which is polymerised to make a specific protein.

Genetic engineering - When the genome of an organism is modified to change its characteristics.

Genetic profiling - The process of creating a 'profile' for an individual based on their DNA. The DNA is cut into bands and separated into a characteristic band pattern, unique to the individual.

Genotype - The genetic makeup of an organism.

Heterozygous - When an individual has two non-identical alleles of a gene e.g. Bb.

Homozygous - When an individual has two identical alleles of a gene e.g. bb.

Meiosis - A form of cell division that produces gametes. They are not genetically identical and contain half the number of chromosomes.

Nucleotide - The monomers of DNA that consist of a common sugar, a phosphate group and a base attached to the sugar. The chemical base can be one of A, C, T or G.

PKU (Phenylketonuria) - A genetic disorder caused by homozygous recessive alleles.

Phenotype - The physical characteristics of an organism. It is due to interactions between the genotype and the environment.

Protein synthesis - The formation of proteins from amino acids which takes place in the ribosomes.

Punnett square - A grid used to determine potential outcomes of a genetic cross.

Recessive - An allele that is only expressed if two copies are present. It is represented by a small letter.

Selfing - Self-fertilisation in plants. Selfing occurs when pollen from a plant lands on the stigma of the same plant.

Sex chromosomes - A pair of chromosomes responsible for the determination of gender. XY in males, XX in females.

Single gene inheritance (monohybrid inheritance) - Inheritance of characteristics that are controlled by a single gene.

Triplet code - The triplet code is a code of three bases. Three bases codes for a specific amino acid.

→ Definition taken from: WJEC (Wales) GCSE in BIOLOGY Specification V.2 January 2019







