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3 Edexcel GCSE Biology



Reproduction

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Types of Reproduction

Your notes

Sexual Reproduction

Sexual reproduction is:

The process involving the fusion of the nuclei of two gametes to form a zygote (fertilised egg cell) and the production of offspring that are genetically different from each other

• Fertilisation is defined as the **fusion of gamete nuclei**, and as each gamete comes from a **different parent**, there is **variation** in the offspring

Gametes and zygotes

- A gamete is a sex cell (in animals: sperm and ovum; in plants: pollen nucleus and ovum)
- Gametes differ from normal cells as they contain half the number of chromosomes found in other body cells - we say they have a haploid nucleus
- This is because they only contain one copy of each chromosome, rather than the two copies found in other body cells
- In human beings, a normal body cell contains 46 chromosomes but each gamete contains 23
 chromosomes
- When the male and female gametes fuse, they become a **zygote** (fertilised egg cell)
- This contains the full **46 chromosomes**, half of which came from the father and half from the mother we say the zygote has a **diploid nucleus**

Advantages and disadvantages of sexual reproduction

• Although there are some clear advantages of sexual reproduction (in comparison to asexual reproduction), there are also a few disadvantages of sexual reproduction

Advantages & Disadvantages of Sexual Reproduction Table



ADVANTAGES	DISADVANTAGES
INCREASES GENETIC VARIATION	TAKES TIME AND ENERGY TO FIND MATES
THE SPECIES CAN ADAPT TO NEW ENVIRONMENTS DUE TO VARIATION, GIVING THEM A SURVIVAL ADVANTAGE	DIFFICULT FOR ISOLATED MEMBERS OF THE SPECIES TO REPRODUCE
DISEASE IS LESS LIKELY TO AFFECT POPULATION (DUE TO VARIATION)	



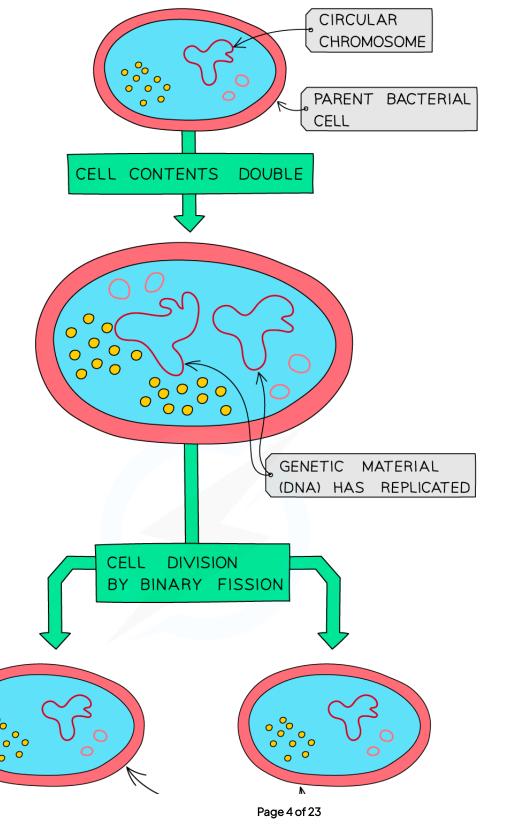
Asexual Reproduction

Asexual reproduction is:

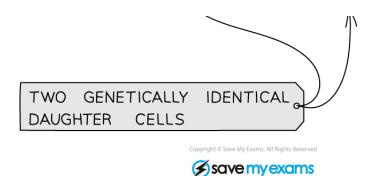
The process resulting in genetically identical offspring being produced from one parent

- Asexual reproduction does not involve gametes or fertilisation
- Only one parent is required so there is no fusion of gametes and no mixing of genetic information
- As a result, the offspring are genetically identical to the parent and to each other (they are clones)
- Many **plants** reproduce via asexual reproduction
- Bacteria produce exact genetic copies of themselves in a type of asexual reproduction called binary fission











Bacteria produce exact genetic copies of themselves in a type of asexual reproduction called binary fission

Advantages and disadvantages of asexual reproduction

 Whilst asexual reproduction has several advantages over sexual reproduction, it also has a few disadvantages

Advantages & Disadvantages of Asexual Reproduction Table

ADVANTAGES	DISADVANTAGES
POPULATION CAN BE INCREASED RAPIDLY WHEN CONDITIONS ARE RIGHT	LIMITED GENETIC VARIATION IN POPULATION - OFFSPRING ARE GENETICALLY IDENTICAL TO THEIR PARENTS
CAN EXPLOIT SUITABLE ENVIRONMENTS QUICKLY	POPULATION IS VULNERABLE TO CHANGES IN CONDITIONS AND MAY ONLY BE SUITED FOR ONE HABITAT
MORE TIME AND ENERGY EFFICIENT	DISEASE IS LIKELY TO AFFECT THE WHOLE POPULATION AS THERE IS NO GENETIC VARIATION
REPRODUCTION IS COMPLETED MUCH FASTER THAN SEXUAL REPRODUCTION	

Comparing sexual and asexual reproduction

- The key differences between sexual and asexual reproduction include:
 - The number of parent organisms
 - How offspring are produced (the type of cell division required)



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- The level of genetic similarity between offspring
- The possible **sources of genetic variation** in offspring
- The **number of offspring** produced
- The **time taken** to produce offspring

Comparing Sexual & Asexual Reproduction Table

Feature	Asexual Reproduction	Sexual Reproduction
Number of parent organisms	One	Two
Type of cell division required	Offspring are produced by mitosis	Offspring are produced from the fusion of two haploid gametes, which are produced by meiosis
Level of genetic similarity between offspring	Offspring are genetically identical to each other (and to the parent)	Offspring are geenetically unique (and genetically different from both parents)
Sources of genetic variation in offspring	Only one source: mutation	Three sources: the contribution of 50% of their DNA from each of the two parents, the production of gametes by meiosis (resulting in new combinations of alleles on the chromosomes), the random distribution of chromosomes into gametes during meiosis
Number of offspring produced	Usually relatively large numbers	Usually relatively limited numbers
Time taken to produce offspring	Usually relatively fast Copyright © Save My Exams, All P	Usually relatively slowly

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The Role of Meiosis

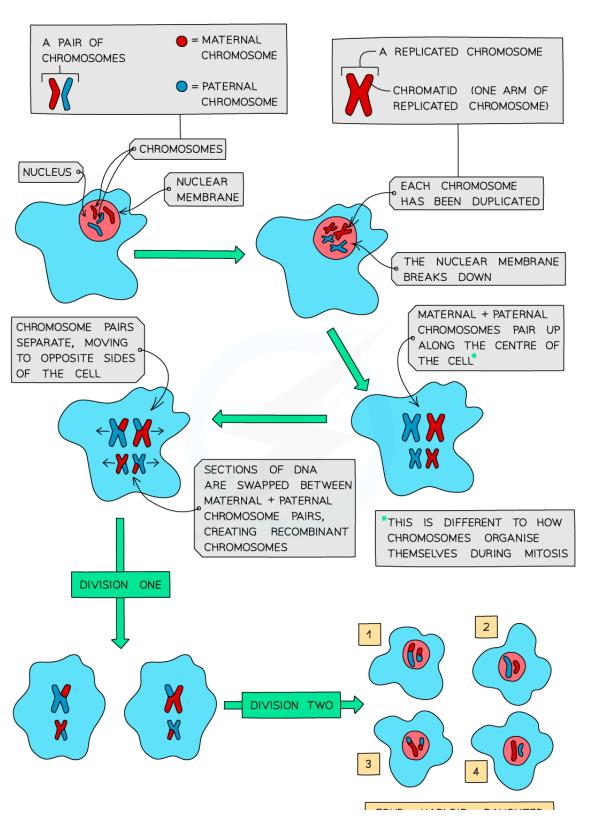
Your notes

The Role of Meiosis

- Cells in **reproductive organs** divide by **meiosis** to form **gametes** (sex cells)
- The number of chromosomes must be halved when the gametes are formed
- Otherwise, there would be double the number of chromosomes after they join at **fertilisation** to form the zygote (fertilized egg)
- This halving occurs during meiosis, and so it is described as a reduction division in which the chromosome number is halved from diploid to haploid, resulting in genetically different haploid gametes
- It starts with chromosomes doubling themselves as in mitosis and lining up in the centre of the cell
- After this has happened the cells divide twice so that only one copy of each chromosome passes to each gamete
- We describe gametes as being haploid having half the normal number of chromosomes
- Because of this double division, meiosis produces four haploid daughter cells



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FOUR HAPLOID DAUGHTER
CELLS PRODUCED

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The importance of meiosis

- Produces gametes e.g. sperm cells and egg cells in animals or pollen grains and ovum cells in plants
- Increases genetic variation of offspring
- Meiosis produces variation by forming new combinations of maternal and paternal chromosomes every time a gamete is made, meaning that when gametes fuse randomly at fertilisation, each offspring will be genetically different from any others



Examiner Tips and Tricks

You are not required to know the **stages** of meiosis for your exam. They are only included here to help with your understanding of the **role** of meiosis (i.e. the production of **four daughter cells**, each with **half** the number of **chromosomes**, which results in **genetically different haploid gametes**).





DNA & the Genome

Your notes

The Genome

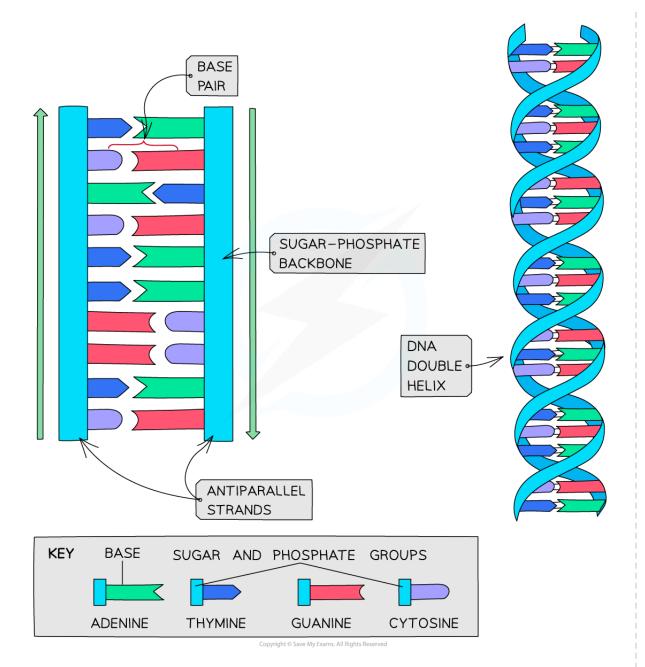
- The entire set of the genetic material of an organism is known as its genome
- In 2003, scientists completed a 13-year project in which they sequenced the **genes** that make up the **whole human genome**
 - This project was named the **human genome project**
- A gene is a section of DNA that codes for a particular sequence of amino acids
- These sequences of amino acids form **proteins**
- Genes control our characteristics as they code for proteins that play important roles in what our cells do

The Structure of DNA

- DNA, or deoxyribonucleic acid, is the genetic material found in the nucleus of a cell
- DNA is a **polymer** made up of **two strands** coiled around to make a **double helix**



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The structure of DNA and complementary base pairs

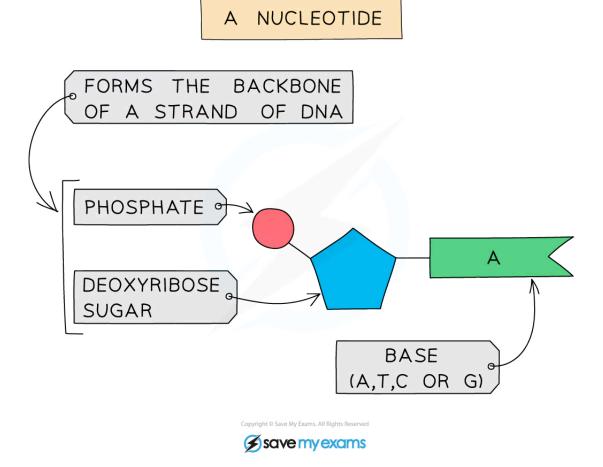
Nucleotides

- DNA is a **polymer** (a molecule made from many **repeating subunits**)
- These individual subunits of DNA are called **nucleotides**



 Each nucleotide consists of a common sugar and phosphate group with one of four different bases attached to the sugar

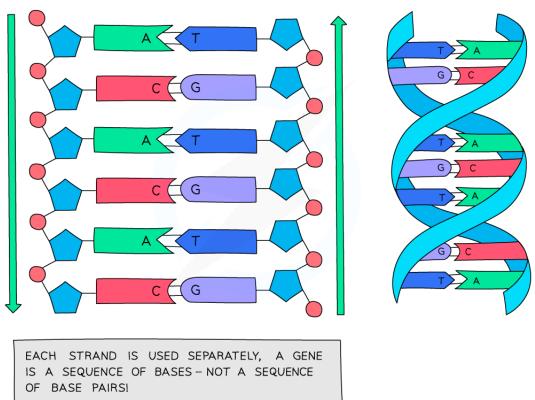




A nucleotide

• The **phosphate and sugar section of the nucleotides** form the **'backbone'** of the DNA strand (like the sides of a ladder) and the base pairs of each strand connect to form the rungs of the ladder







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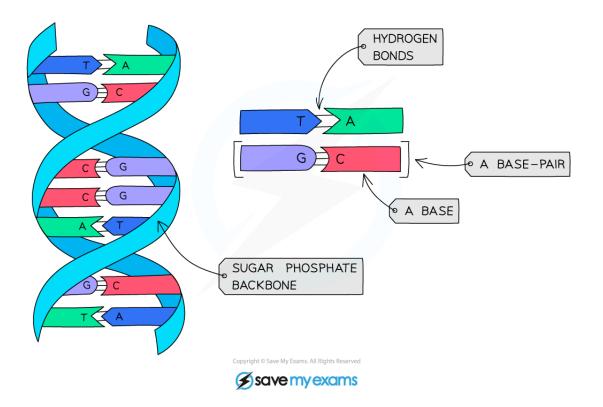
DNA is a polynucleotide - this means it is a polymer made up of many repeating subunits (monomers) known as nucleotides

Base pairing

- There are four different nucleotides
- These four nucleotides contain the same phosphate and deoxyribose sugar, but differ from each other in the base attached
- There are **four different bases**: Adenine (**A**), Cytosine (**C**), Thymine (**T**) and Guanine (**G**)
- The bases on each strand pair up with each other, holding the two strands of DNA in the double helix together
- The bases always pair up in the same way:
 - Adenine always pairs with Thymine (A-T)



- Cytosine always pairs with Guanine (C-G)
- This is known as 'complementary base pairing'
- The **complementary base pairs** (A-T and C-G) pair up by forming **weak hydrogen bonds** with each other





Extracting DNA

- DNA can actually be **extracted from fruit** using some basic classroom equipment
- Fruits that have relatively large amounts of DNA in their cells, such as strawberries, bananas and kiwis, can be used

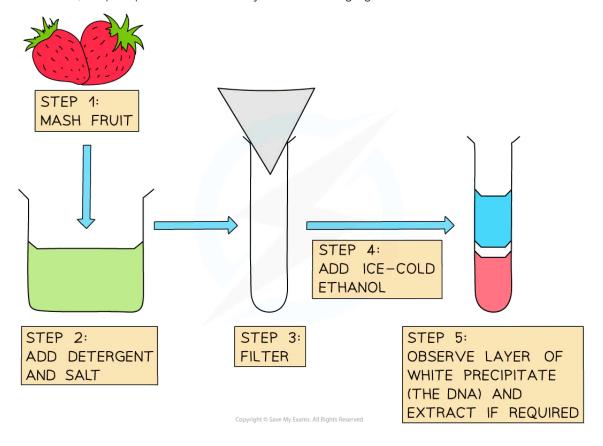
Method

- Mash the fruit and mix this into a beaker containing a solution of detergent (e.g. washing up liquid) and salt
 - The detergent **breaks down** the **cell membranes** (and the **nuclear membranes**), causing the fruit cells to **release their DNA**





- The salt causes the DNA to stick together
- Filter the mixture into a test tube
 - This **removes any debris** (big, insoluble bits of cell) from the mixture
- Gently add some ice-cold ethanol to the filtrate (the filtered mixture) by pouring the ethanol slowly
 down the side of the test tube
 - This causes the DNA to precipitate (i.e. to come out of solution) as DNA is not soluble in cold alcohol
- The DNA will appear as a **stringy white precipitate** (a solid)
- If needed, this precipitate can be carefully **extracted** using a glass rod



A simple method for extracting DNA from fruit





Protein Synthesis

Your notes

Genes

Higher tier only

- A gene is a section of DNA with a particular sequence of bases that codes for a particular sequence of amino acids
- A sequence of **three bases** is the **code** for a single specific **amino acid**
- The order of bases controls the order and different types of amino acids that are eventually joined together in a polypeptide chain
- These polypeptide chains (the amino acid sequences) then form a particular type of **protein**
- In this way, it is the order of bases in the DNA that eventually determines which proteins are produced

Types of proteins

- When the protein chain is complete it folds up to form a unique shape
- This unique shape enables the proteins to fulfil a specific function. For example, proteins can be:
 - Enzymes proteins that act as biological catalysts to speed up chemical reactions occurring in the body (e.g. maltase is an enzyme that breaks down maltose into glucose)
 - Hormones proteins that carry messages around the body (e.g. testosterone is a hormone that
 plays an important role in the development of the male reproductive system and development of
 male secondary sexual characteristics, such as increased muscle mass and growth of body hair)
 - Structural proteins proteins that provide structure and are physically strong (e.g. collagen is a structural protein that strengthens connective tissues such as ligaments and cartilage)
- There are also many other types of proteins

Transcription & Translation

Higher tier only

- A gene is a sequence of nucleotide bases in a DNA molecule that codes for the production of a specific sequence of amino acids, that in turn make up a protein
- This process of protein synthesis occurs in **two stages**:
 - Transcription DNA is transcribed and an mRNA molecule is produced
 - Translation mRNA (messenger RNA) is translated and an amino acid sequence is produced



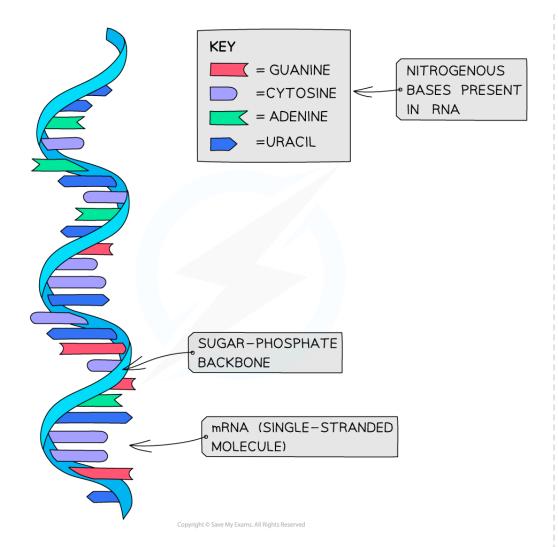
Transcription

- This stage of protein synthesis occurs in the **nucleus** of the cell
- Part of a DNA molecule **unwinds** (the hydrogen bonds between the complementary base pairs break)
- This exposes the gene to be transcribed (the gene from which a particular protein will be produced)
- An enzyme known as RNA polymerase binds to a region of non-coding DNA just in front of the gene
- RNA polymerase then makes a complementary copy of the code from the gene by building a singlestranded nucleic acid molecule known as mRNA (messenger RNA)
- The mRNA molecule **leaves the nucleus** via a pore in the nuclear envelope
 - Like DNA, the nucleic acid RNA (ribonucleic acid) is a polynucleotide it is made up of many nucleotides linked together in a long chain
 - Like DNA, RNA nucleotides contain the nitrogenous bases adenine (A), guanine (G) and cytosine
 (C)
 - **Unlike DNA**, RNA nucleotides never contain the nitrogenous base thymine (T) in place of this they contain the nitrogenous base **uracil (U)**
 - Unlike DNA, RNA molecules are only made up of one polynucleotide strand (they are singlestranded)
 - Each RNA polynucleotide strand is made up of alternating ribose sugars and phosphate groups linked together, with the nitrogenous bases of each nucleotide projecting out sideways from the single-stranded RNA molecule
 - One example of an RNA molecule is messenger RNA (mRNA), which is the transcript copy of a gene
 that encodes a specific polypeptide. Two other examples are transfer RNA (tRNA) and ribosomal
 RNA (rRNA)





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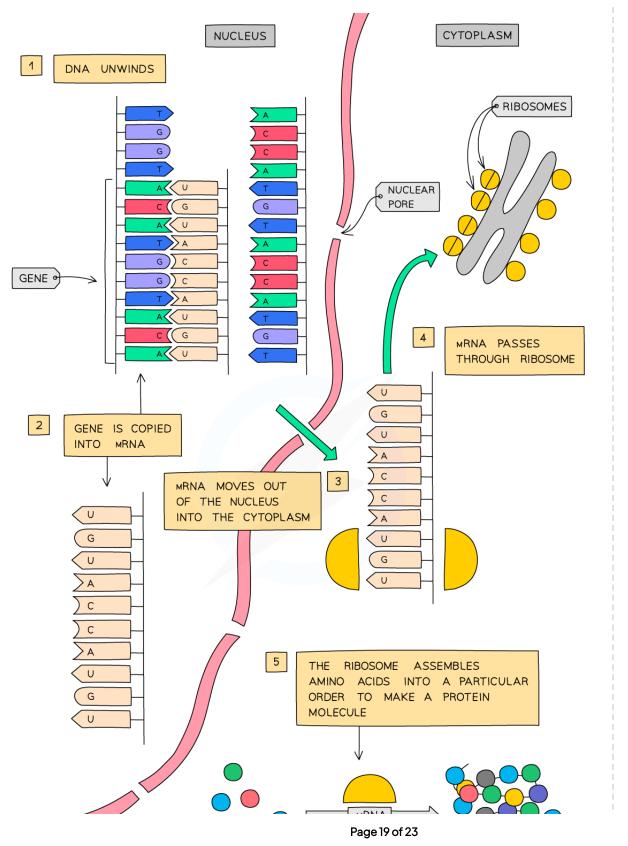




Messenger RNA (mRNA) provides a good example of the structure of RNA

Translation

- This stage of protein synthesis occurs in the **cytoplasm** of the cell
- After leaving the nucleus, the mRNA molecule attaches to a **ribosome**
 - The ribosome 'reads' the code on the mRNA in groups of three
 - Each **triplet of bases** on the mRNA molecule (known as a codon) codes for a **specific amino acid**
 - In this way, the ribosome **translates** the sequence of **bases** into a sequence of **amino acids** that make up a **protein**

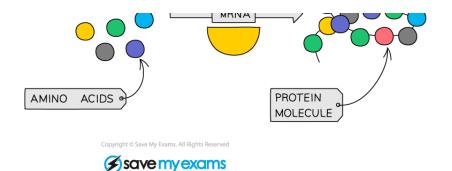




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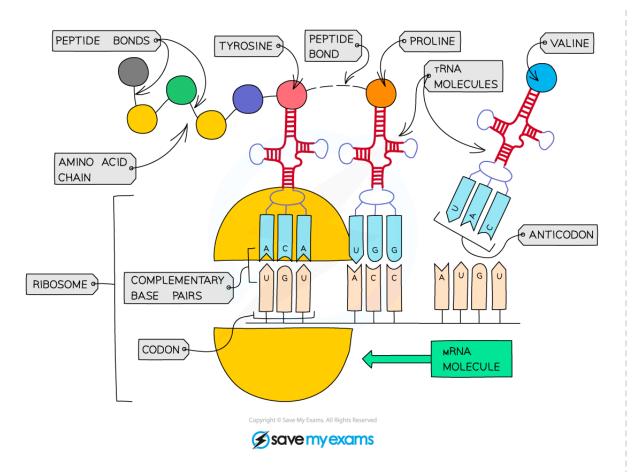
An overview of protein synthesis

- In the cytoplasm, there are free molecules of **tRNA** (transfer RNA)
- These tRNA molecules have a triplet of unpaired bases at one end (known as the anticodon) and a region where a specific amino acid can attach at the other
 - There are at least 20 different tRNA molecules, each with a specific anticodon and specific amino acid binding site
- The tRNA molecules bind with their specific amino acids (also in the cytoplasm) and bring them to the mRNA molecule on the ribosome
- The triplet of bases (anticodon) on each tRNA molecule pairs with a complementary triplet (codon) on the mRNA molecule
- Two tRNA molecules fit onto the ribosome at any one time, bringing the amino acid they are each carrying side by side
- A **peptide bond** is then formed between the two amino acids
- This process continues until a 'stop' codon on the mRNA molecule is reached this acts as a signal for translation to stop and at this point the amino acid chain coded for by the mRNA molecule is complete
- This amino acid chain (the final polypeptide) will then fold to form a protein



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The triplet of bases (codon) of mRNA is read by the ribosome and amino acids are attached together in a specific sequence to form the protein



Examiner Tips and Tricks

In an exam, you could be asked why DNA must first be transcribed into mRNA before it can be translated into a protein. This is necessary because **DNA cannot travel out of the nucleus to the ribosomes** (it is far **too big** to pass through a nuclear pore) so the base code of each gene is transcribed into an RNA molecule called messenger RNA (mRNA).

Genetic Variants

Your notes

Genetic Variants in Non-coding DNA

Higher tier only

- During transcription, part of a DNA molecule unwinds, exposing the gene to be transcribed (the gene from which a particular protein will be produced)
- An enzyme known as RNA polymerase then binds to a region of non-coding DNA just in front of this
 gene
 - Non-coding means that the DNA bases do not code for the production of an amino acid
 - RNA polymerase makes a complimentary copy of the gene by building a single-stranded nucleic acid molecule known as mRNA (messenger RNA)
- However, genetic variants (different versions of genes caused by mutations) can be produced if a mutation occurs in the non-coding DNA of a gene
 - A mutation is a random change to the DNA base sequence
- Such genetic variants can affect phenotype by influencing the binding of RNA polymerase and altering the quantity of protein produced:
 - If a **mutation** happens in the non-coding region to which RNA polymerase attaches, this can affect the **ability** of RNA polymerase to **bind** to it
 - In some cases, this might make it easier for RNA polymerase to bind. In other cases, it might make it more difficult
 - If it makes it more difficult for RNA polymerase to bind, less mRNA will be transcribed from the gene and therefore less protein (that the gene codes for) will be produced
 - Depending on the function of this protein, the phenotype of the organism may be affected by how much of it is produced
 - In this way, genetic variants in the non-coding DNA of a gene can affect phenotype, even if the coding region of the gene is completely normal

Genetic Variants in Coding DNA

Higher tier only

 Genetic variants (different versions of genes caused by mutations) can also be produced if a mutation occurs in the coding DNA of a gene



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- If a mutation happens in the coding region of a gene, the gene may code for a **different sequence of** amino acids
- Your notes
- As the amino acid sequences are highly specific and form specific shapes and types of proteins, a
 change in the sequence may change the shape of the final protein and therefore may affect how it
 functions
 - For example, if its amino acid sequence is changed, the activity of a protein such as an enzyme may be increased, decreased or stopped altogether
- This, in turn, could change the **phenotype** (the **characteristics**) of the organism
 - For example, XDH is an enzyme found in fruit flies. Fruit flies with normal genes coding for XDH show normal levels of XDH activity and have red eyes. Fruit flies with genetic variants of the genes coding for XDH (caused by mutations in the coding DNA) show decreased levels of XDH activity and have brown eyes, as they cannot produce the red eye pigment