



The general structure of a DNA nucleotide. An RNA nucleotide has an additional oxygen molecule in the five-carbon sugar ring. Notice the numbering of the carbon atoms on the sugar molecule. The five carbon atoms of the sugar of the nucleotide are numbered 1' to 5', and they proceed clockwise from the oxygen atom. The prime symbol (') indicates that the carbon belongs to the sugar rather than to the base.



Nucleotides are joined together in a long chain. The "backbone" of the chain is made up of alternating sugar and phosphate groups that are joined by chemical bonds. The nitrogenous bases project out from the sugar-phosphate backbone.



A DNA molecule is made up of two strands of nucleotides that are wound around each other (A). The two strands are held together by hydrogen bonds between complementary base pairs. C-G pairs are held together by three hydrogen bonds, and A-T pairs are held together by two hydrogen bonds. Notice that the chains are antiparallel—the 5' to 3' orientation runs in the opposite direction on each strand. Another example of antiparallelism is represented by M.C. Escher's sketch, "Drawing Hands" (B).



Replication takes place at several locations simultaneously. Each replication bubble represents two replication forks moving in opposite directions along the length of the chromosome. As replication proceeds along the strand, the bubbles grow until they meet. The parent strand of DNA is shown in grey. The new complementary strand is shown in black.



During DNA synthesis, the overall direction of elongation is the same along both strands, but elongation occurs differently. On the leading strand, DNA synthesis takes place along the DNA molecule in the same direction as the movement of the replication fork. On the lagging strand, DNA synthesis proceeds in the opposite direction to the movement of the replication fork. As well, the lagging strand is synthesized in short fragments.



This simplified illustration of the replication machine shows how a loop in the lagging strand allows a single polymerase complex to replicate both DNA strands simultaneously.



The path of gene expression. The "central dogma" proposes that genetic information passes (via transcription) from the genes (DNA) to an RNA copy of the gene, and the RNA copy directs the sequential assembly of a chain of amino acids to produce a protein (via translation).

mRNA

<u>mRNA</u>

- Ribonucleic acid
- Sugar component is ribose
- Single stranded; may fold in on itself to form regions of complementary base pairs
- Nitrogenous bases are A, U, C, and G
- One of three forms of RNA: mRNA, tRNA, and rRNA
- Carries genetic information from DNA to ribosomes for translation
- In eukaryotes, mRNA is found in the mitochondria, nucleus, and cytoplasm

DNA

- Deoxyribonucleic acid
- Sugar component is deoxyribose
- Double stranded; double helix
- Nitrogenous bases are A, T, C, and G
- Only one form of DNA
- Stores genetic information that is replicated during cell division and copied into mRNA during transcription
- In eukaryotes, DNA is found in the nucleus and mitochondria

CHAPTER 18

OVERHEAD

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Table: Messenger RNA Codons and Their Corresponding Amino Acids

First base	Second base				Third base
	U	С	Α	G	
U	UUU phenylalanine	UCU serine	UAU tyrosine	UGU cysteine	U
	UUC phenylalanine	UCC serine	UAC tyrosine	UGC cysteine	С
	UUA leucine	UCA serine	UAA stop**	UGA stop**	Α
	UUG leucine	UCG serine	UAG stop**	UGG tryptophan	G
С	CUU leucine	CCU proline	CAU histidine	CGU arginine	U
	CUC leucine	CCC proline	CAC histidine	CGC arginine	С
	CUA leucine	CCA proline	CAA glutamine	CGA arginine	Α
	CUG leucine	CCG proline	CAG glutamine	CGG arginine	G
Α	AUU isoleucine	ACU threonine	AAU asparagine	AGU serine	U
	AUC isoleucine	ACC threonine	AAC asparagine	AGC serine	С
	AUA isoleucine	ACA threonine	AAA lysine	AGA arginine	Α
	AUG methionine*	ACG threonine	AAG lysine	AGG arginine	G
G	GUU valine	GCU alanine	GAU aspartate	GGU glycine	U
	GUC valine	GCC alanine	GAC aspartate	GGC glycine	C
	GUA valine	GCA alanine	GAA glutamate	GGA glycine	A
	GUG valine	GCG alanine	GAG glutamate	GGG glycine	G

* AUG is an initiator codon. It also codes for the amino acid methionine.

** UAA, UAG, and UGA are terminator codons.



During transcription, a complex of RNA polymerases track along the DNA molecule, synthesizing a single-stranded mRNA molecule that is complementary to the sense strand of DNA. The DNA helix reforms behind the RNA polymerase complex.





Each tRNA molecule is about 80 nucleotides long. The tRNA molecule shown here has the anticodon GCU, which pairs with the mRNA sequence CGA. This tRNA molecule carries the amino acid arginine.



The translation cycle involves the base pairing of a new tRNA, the transfer of a growing polypeptide chain, and the movement of the ribosome along the mRNA molecule. The complete cycle occurs about 15 times per second in a prokaryotic cell.



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Gene Expression

Role of Various Nucleic Acids in Gene Expression

Nucleic Acid	Structure	Function
DNA	Double helix	Stores genetic information that is copied
		into mRNA during transcription
mRNA	Linear single	Carries genetic information from DNA
	strand	to the ribosomes; in eukaryotes, mRNA
		is processed before it moves to the
		cytoplasm for translation
tRNA	Lobed shape	Carries a particular amino acid to the
		correct mRNA codon site during
		translation
rRNA	Linear single	Combines with a complex of proteins to
	strand	form ribosomes, which bring together
		the mRNA strand and tRNA molecules
		during translation

premature stop codon. No functional polypeptide will be produced from

this gene.

CHAPTER 18 BLM 18.3.1 **Mutations OVERHEAD** GUU-CAU-UUG-ACU-CCC-GAA-GAA val – his – leu – thr – pro – glu – glu A The normal coding sequence, with the codons in the top row and the resulting amino acids below them. GUU-CAU-UUG-ACU-CCC-GAA-GAA val – his – leu – thr – pro – glu – glu GUU-CAU-UUG-ACC-CCC-GAA-GAA val – his – leu – thr – pro – glu – glu A The normal coding sequence, with the codons in the top row and the B This mutation is silent, since the resulting amino acids below them. change in nucleotide sequence has no effect on the polypeptide product. GUU-CAU-GUU-GAC-UCC-CGA-AGA A GUU-CAU-UUG-ACU-CCC-GUA-GAA val – his – val – asp – ser – arg – arg val – his – leu – thr – pro – val – glu B The insertion of a single nucleotide, C This is a mis-sense mutation, since it in this case quanine, results in a causes the amino acid valine to be frameshift mutation. inserted in the place of glutamate within the polypeptide chain. The resulting protein is unable to transport oxygen effectively and produces a GUU-CAU-UUG-CUC-CCG-AAG-AA disorder known as sickle cell disease. val – his – leu – leu – pro – lys C Similarly, a deletion of even a single GUU-CAU-UAG nucleotide, in this case adenine, also val – his – stop results in a frameshift mutation. This substitution causes a nonsense mutation by changing the codon for the amino acid leucine (UUG) into a

(Left) A nucleotide substitution can have varied effects, as shown on this portion of the gene that codes for human betaglobulin, one of the two polypeptides in the blood protein hemoglobin. (Right) Frameshift mutations are usually nonsense mutations.



Gel Electrophoresis

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- A Restriction enzymes Either one or several restriction enzymes are added to a sample of DNA. The enzymes cut the DNA into fragments.





Before the DNA fragments are added to the wells, they are treated with a dye that glows under ultraviolet light, allowing the bands to be studied.



B The gel A gel, with a consistency similar to gelatin, is formed so small wells are left at one end. Small amounts of the DNA sample are placed into these wells.



C The electrical field The gel is placed in a solution, and an electrical field is set up so one end of the gel is positive and the other end is negative.

D The fragments move The negatively charged DNA fragments travel toward the positive end. The smaller the fragment, the faster it moves through the gel. Fragments that are the farthest from the well are the smallest.