

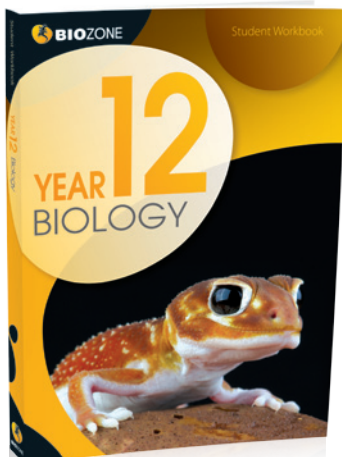
# 12

YEAR

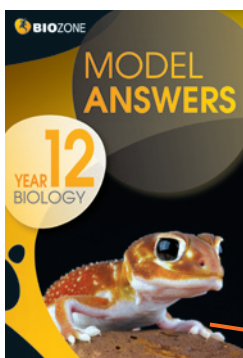
# BIOLOGY



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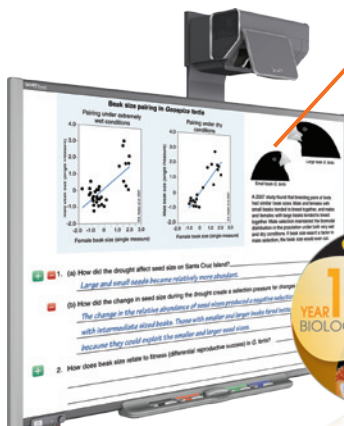
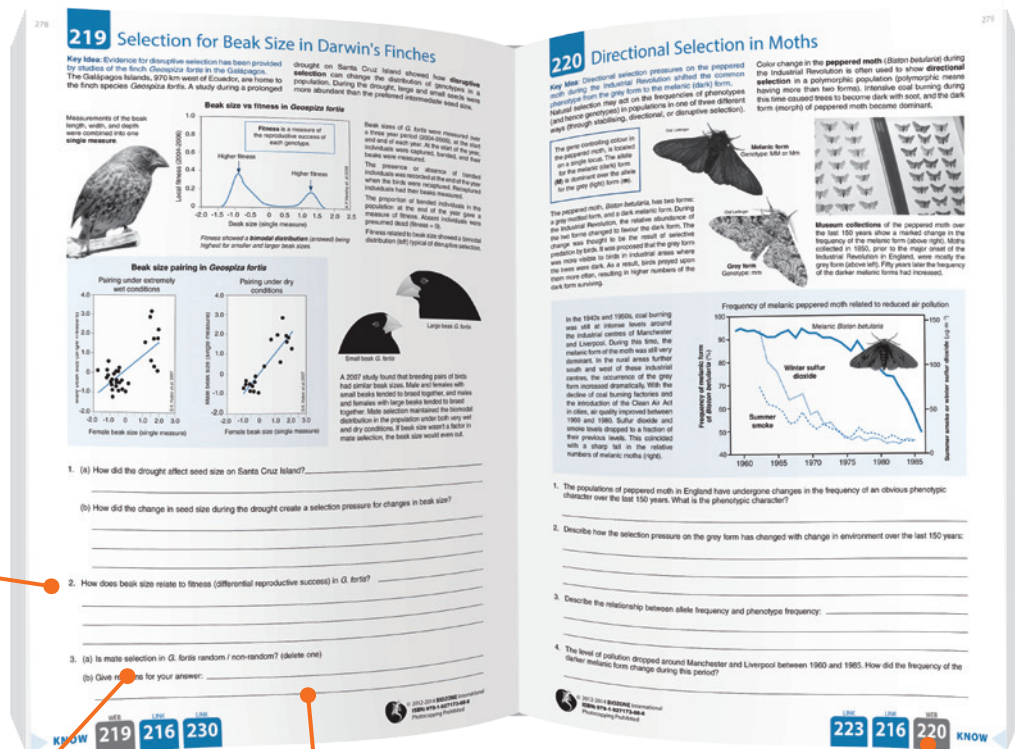
**Engage** students with write-on activities directly in the workbook



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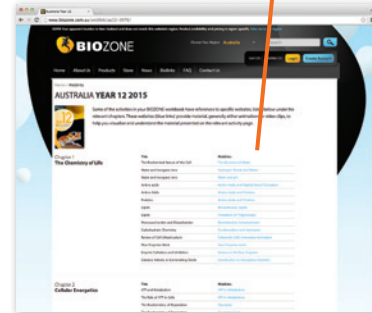
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## SIGNIFICANT BENEFITS

**Streamline** the coverage of extensive curricula.

**Concept-based** content allows complex ideas to be broken down into manageable components.

**Student engagement** High level of student engagement with information, questions, and the student's answers all on the same page.

**Critical-thinking** questions challenge student understanding - great for test preparation.

**Visually rich** content, including clear explanatory diagrams, appeals to today's learning styles.



# Defence and the Immune System

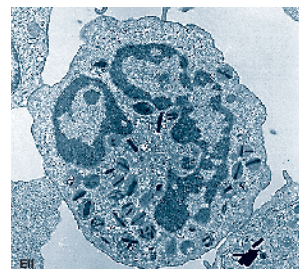
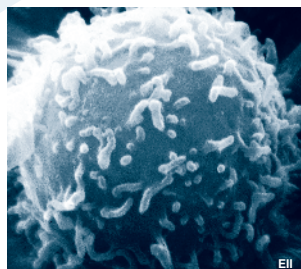
## Key terms

active immunity  
allergic reaction  
antibody  
(=immunoglobulin)  
antigen  
asthma  
autoimmune disease  
B cell (=B lymphocyte)  
cell-mediated immunity  
clonal selection  
fever  
hayfever  
histamine  
humoral immunity  
hypersensitivity  
immunity  
immunological memory  
infection  
inflammation  
interferon  
leucocyte  
lymphocyte  
macrophage  
MHC  
monoclonal antibody  
non-specific defences  
(=innate immunity)  
passive immunity  
phagocyte  
primary response  
secondary response  
specific (=adaptive)  
immune response  
T cell (=T lymphocyte)  
thymus  
vaccination  
(=immunisation)

## The body's layers of defence

### Learning aims and skills

- ☐ 1 Describe the range of physical and chemical defences in animals. **91**
- ☐ 2 Explain how the body distinguishes self from non-self. Include reference to the role of the major histocompatibility complex (MHC) and its role in self-recognition and in determining tissue compatibility in transplant recipients. **92**
- ☐ 3 Explain the basis of the Rh and ABO blood group systems in humans. Explain the consequences of blood type incompatibility in blood transfusions. **93**
- ☐ 4 Describe non-specific (innate) defences in humans, describing the nature and role of each of the following in protecting against pathogens: **94-97**
  - (a) Skin (including sweat and sebum production) and mucous membranes.
  - (b) Body secretions (tears, urine, saliva, gastric juice).
  - (c) Natural anti-bacterial and anti-viral proteins, e.g. interferon.
  - (d) Phagocytosis, the inflammatory response, fever, and cell death.



Activity number

## Specific immunity

### Learning aims and skills

- ☐ 5 Describe the specific immune response, including the role of specificity and memory. Describe cell-mediated immunity and humoral (antibody-mediated) immunity, identifying the specific white blood cells involved in each case. **98 99**
- ☐ 6 Describe clonal selection and the basis of immunological memory. Explain how the immune system is able to respond to the large range of potential antigens. **100**
- ☐ 7 Explain antibody production, including how B cells bring about humoral (antibody-mediated) immunity to specific antigens. **101**
- ☐ 8 Distinguish between naturally acquired and artificially acquired immunity and between active and passive immunity. **102**
- ☐ 9 Explain the principles of vaccination, including reference to the primary and secondary response to infection and the role of these. **103**
- ☐ 10 Describe the production and applications of monoclonal antibodies. **104 105**

Activity number

## Immune dysfunction and disease

### Learning aims and skills

- ☐ 11 With reference to specific examples, describe the result of immune system dysfunction, e.g. in hypersensitivity reactions such as asthma and hayfever, and in autoimmune diseases such as multiple sclerosis. **106 107**

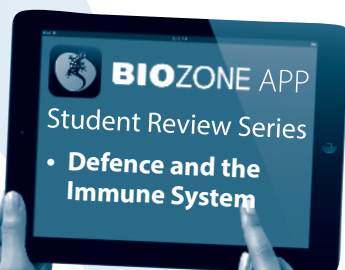
Activity number

## Defence mechanisms in plants

### Learning aims and skills

- ☐ 12 Describe how plants inhibit disease and recover from infection. Describe passive and active defences in plants, including reference to physical and chemical barriers, cellular defences, and cell death (necrosis). **108**

Activity number

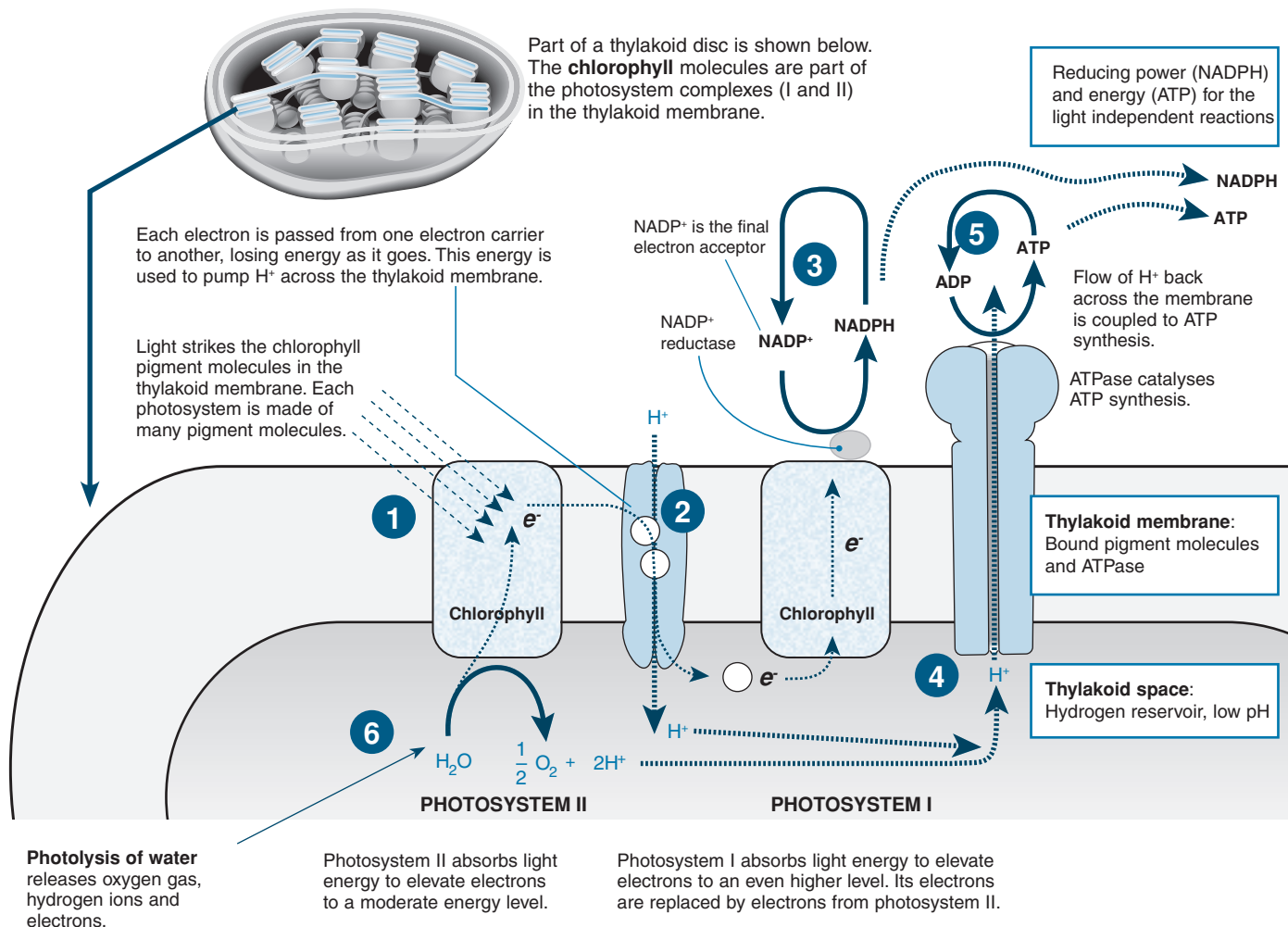


# 35 Light Dependent Reactions

**Key Idea:** In the light dependent reactions of photosynthesis, the energy from photons of light is used to drive the reduction of  $\text{NADP}^+$  and the production of ATP.

The process of photosynthesis occurs in two separate reaction systems. The light dependent reactions (the first of

these) in the thylakoid of the chloroplasts capture light energy to produce **ATP** and reducing power (as **NADPH**) which is used in the light independent reactions (the second reaction system). The light dependent reactions take place in the thylakoids of the chloroplast (enlarged below).



- Where do the light dependent reactions occur? \_\_\_\_\_
- What is the role of each of the following in the light dependent reactions:
  - $\text{NADP}^+$  reductase: \_\_\_\_\_
  - ATPase (also called ATP synthase): \_\_\_\_\_
- How are the electrons that are lost from the thylakoid system replaced? \_\_\_\_\_
- Summarise the events of the light dependent reactions: \_\_\_\_\_

# 55 Neurones and Neurotransmitters

**Key Idea:** Neurones are electrically excitable cells that are specialised to process and transmit information via electrical and chemical signals. Neurotransmitters are chemicals that allow the transmission of signals between neurones.

**Neurones** are cells specialised to transmit information in the

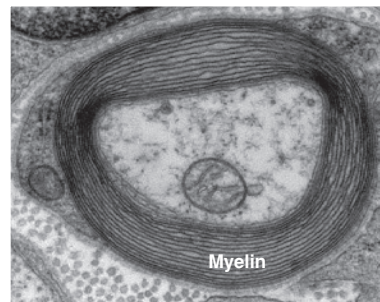
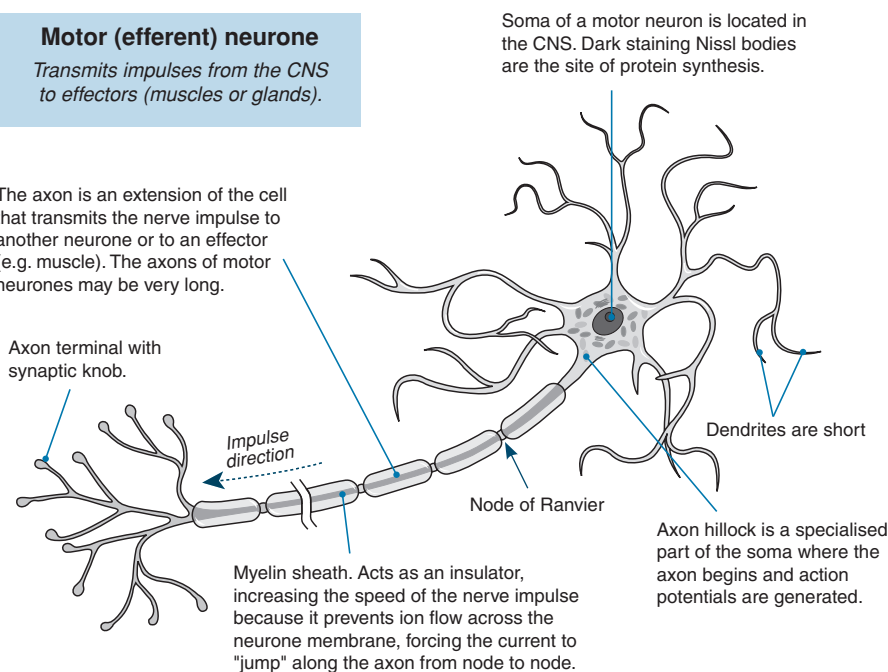
form of electrochemical signals from receptors (in the central nervous system) to effectors. Neurones consist of a cell body (soma) and long processes (dendrites and axons). Messages are transmitted between neurones by signalling molecules called neurotransmitters.

## Motor (efferent) neurone

*Transmits impulses from the CNS to effectors (muscles or glands).*

The axon is an extension of the cell that transmits the nerve impulse to another neurone or to an effector (e.g. muscle). The axons of motor neurones may be very long.

Axon terminal with synaptic knob.

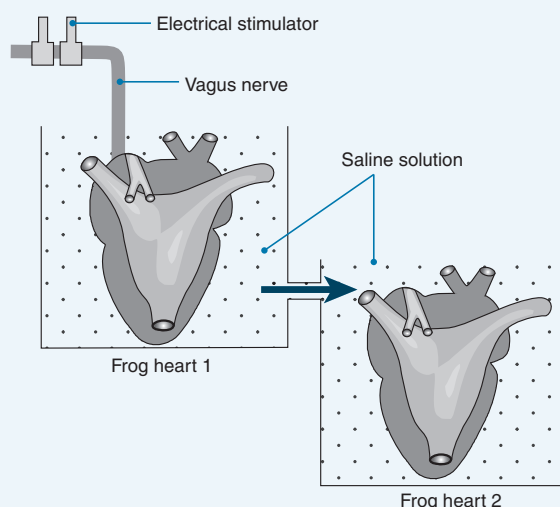
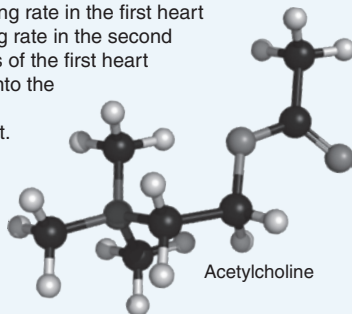


TEM cross section through a myelinated axon

Where conduction speed is important, the axons of neurones are sheathed within a lipid and protein rich substance called **myelin**. Myelin is produced by oligodendrocytes in the central nervous system (CNS) and by Schwann cells in the peripheral nervous system (PNS). At intervals along the axons of myelinated neurones, there are gaps between neighbouring Schwann cells and their sheaths. These are called **nodes of Ranvier**.

## Neurotransmitters Carry Signals Between Neurones

Chemical signalling between neurones was first demonstrated in 1921 by Otto Loewi. In his experiment, the still beating hearts of two frogs were placed in connected flasks filled with saline solution. The vagus nerve (parasympathetic) of the first heart was still attached and was stimulated by electricity to reduce its rate of beating. After a delay, the rate of beating in the second heart also slowed. Increasing the beating rate in the first heart caused an increase in the beating rate in the second heart, showing electrical stimulus of the first heart caused it to release a chemical into the saline solution that then affected the heartbeat of the second heart. The chemical was found to be **acetylcholine**.



1. What is the function of a neurone? \_\_\_\_\_
2. Describe the purpose of a neurotransmitter: \_\_\_\_\_
- 3 (a) Explain why stimulating the first frog heart with electricity caused it to change its beating rate: \_\_\_\_\_
- (b) Explain why the second heart in the experiment reduced its beating rate after a delay: \_\_\_\_\_





# 91 Chemical Defences In Animals

**Key Idea:** Animals have evolved a wide range of chemical defences that protect against infection by pathogens.

Living organisms are under constant attack from many different **pathogens** (disease causing organisms). As a result, they have evolved many ways to limit infection by pathogens.

## Vertebrate Defences

### Antimicrobial substances:

Chemicals (e.g. **lactoferrin**) are secreted from the skin and other body fluids, and also by some white blood cells. These kill pathogens or inhibit their growth.

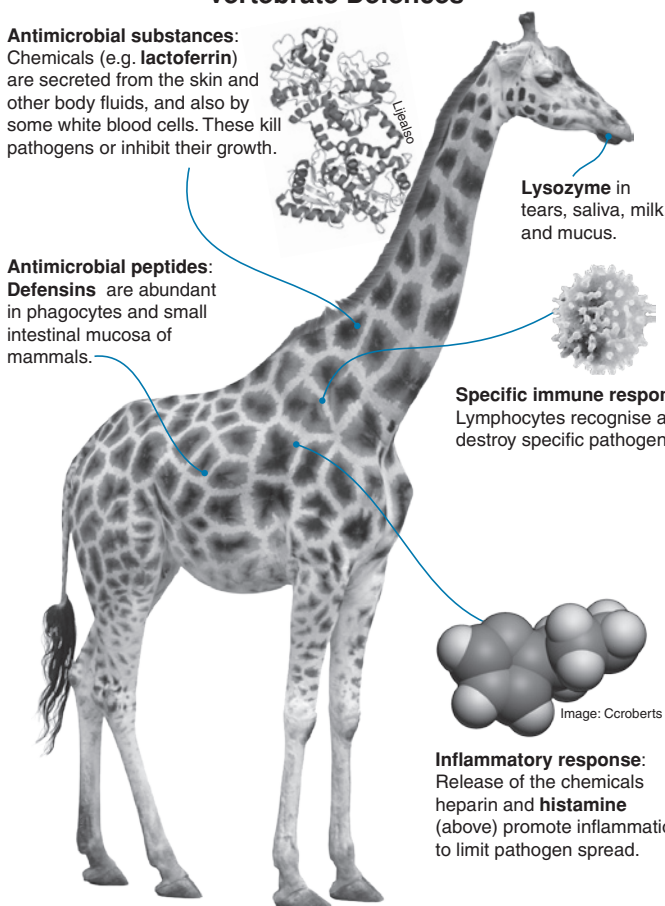
### Antimicrobial peptides:

**Defensins** are abundant in phagocytes and small intestinal mucosa of mammals.

**Lysozyme** in tears, saliva, milk, and mucus.

**Specific immune response:** Lymphocytes recognise and destroy specific pathogens.

**Inflammatory response:** Release of the chemicals **heparin** and **histamine** (above) promote inflammation to limit pathogen spread.



## Invertebrate Defences

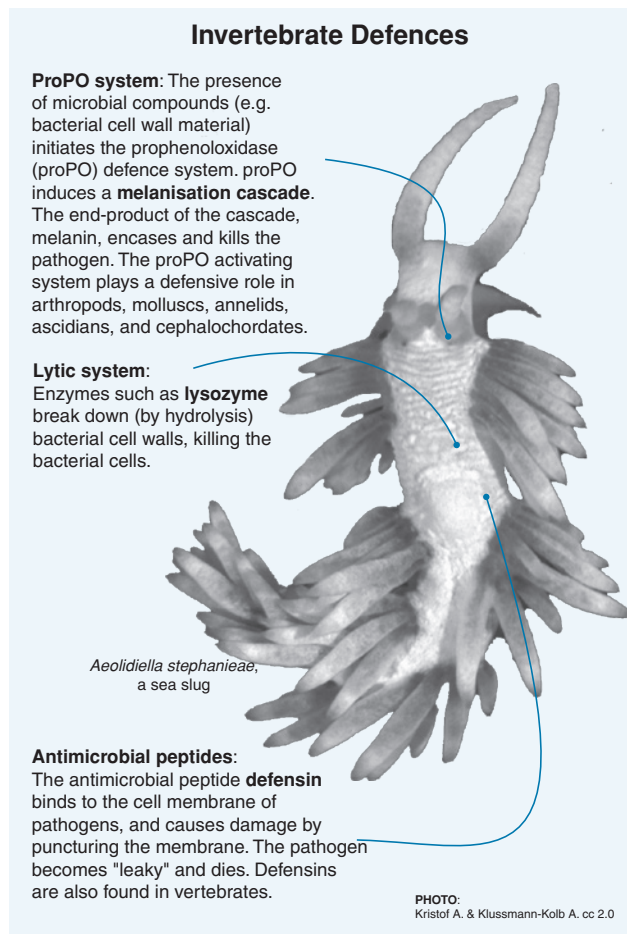
**ProPO system:** The presence of microbial compounds (e.g. bacterial cell wall material) initiates the prophenoloxidase (proPO) defence system. proPO induces a **melanisation cascade**. The end-product of the cascade, melanin, encases and kills the pathogen. The proPO activating system plays a defensive role in arthropods, molluscs, annelids, ascidians, and cephalochordates.

**Lytic system:** Enzymes such as **lysozyme** break down (by hydrolysis) bacterial cell walls, killing the bacterial cells.

**Antimicrobial peptides:** The antimicrobial peptide **defensin** binds to the cell membrane of pathogens, and causes damage by puncturing the membrane. The pathogen becomes "leaky" and dies. Defensins are also found in vertebrates.

*Aeolidiella stephanieae*, a sea slug

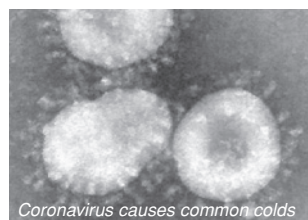
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Kristof A. & Klusmann-Kolb A. cc 2.0



*E. coli* can contaminate water. Pathogenic **bacteria** cause a wide range of diseases. Some types of *E. coli* are harmless gut residents, but others can cause disease.



Mites attached to harvestman host. Vertebrates and invertebrates are host to a range of arthropod vectors, which may carry pathogenic microorganisms on their mouthparts.



Coronavirus causes common colds. **Viruses** are species-specific intracellular parasites and cause many common diseases. They infect all other organisms, including bacteria.



Malaria sporozoite. **Eukaryotic pathogens:** protozoa (e.g. the malaria parasite above), algae, fungi, and parasitic worms, may be pathogenic.

1. (a) Describe the advantage of having multiple (non-specific) defence responses: \_\_\_\_\_

\_\_\_\_\_

(b) Describe a disadvantage of having only general (non-specific) defence responses: \_\_\_\_\_

\_\_\_\_\_

2. Compare and contrast the non-specific defences of vertebrate and invertebrate animals: \_\_\_\_\_

\_\_\_\_\_

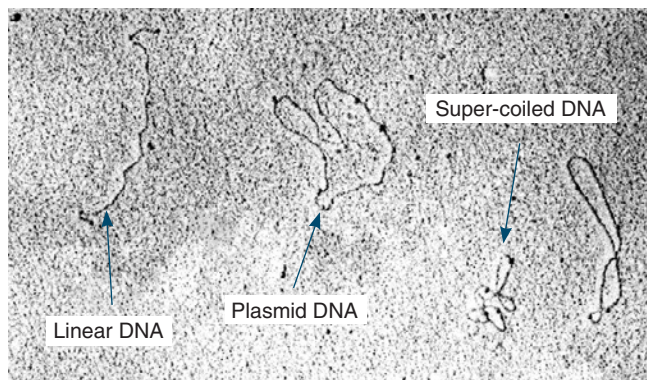
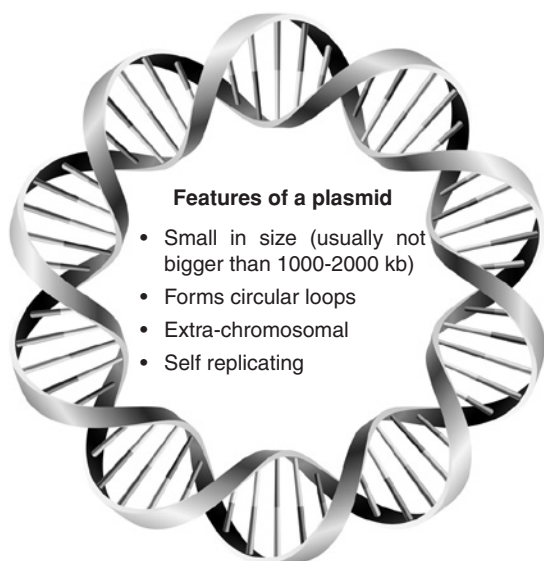
\_\_\_\_\_

# 113 Plasmid DNA

**Key Idea:** Plasmids are small circular pieces of DNA that are not associated with the chromosomal DNA.

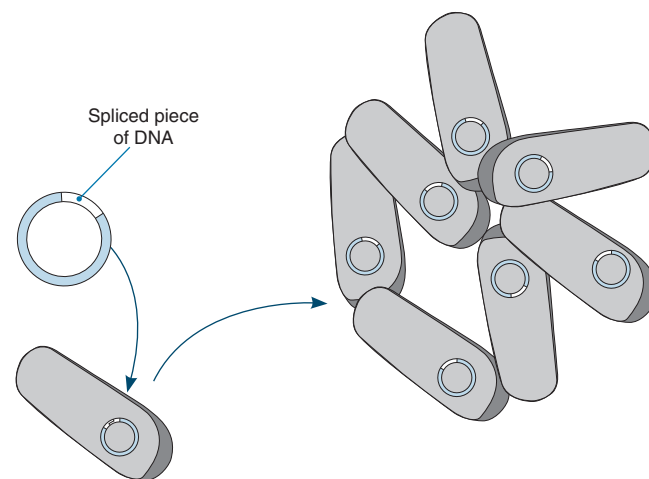
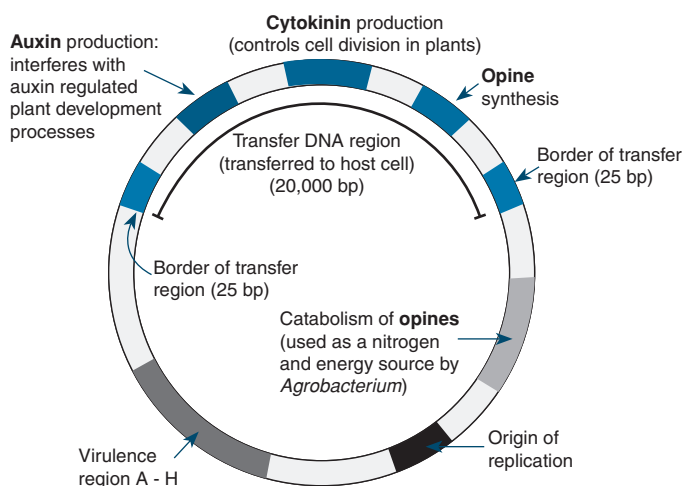
Prokaryotes store most of their genetic information in one large chromosome. However, a small percentage can be found as independently replicating, circular, extra-chromosomal pieces of DNA known as **plasmids**. Plasmids may carry

important genes, such as those for the production of toxins that eliminate prokaryote competitors. Plasmids are less common in eukaryotes but some species, such as the yeast *Saccharomyces*, do have them. The genetic material from viruses may form plasmid-like structures called episomes once they have infected a cell.



Plasmids vary in size from 1000 base pairs (bp) to hundreds of thousands of base pairs. In bacteria, they play an important role in providing extra genetic material that confers properties such as antibiotic resistance.

Plasmids can be transferred between bacterial cells by a process of plasmid transfer called **conjugation**. Conjugation enables bacteria to obtain genetic material from other individuals by **horizontal gene transfer** (transfer of genetic material to individuals other than offspring) and it is an important mechanism for the spread of antibiotic resistance.



The bacterium *Agrobacterium tumefaciens* often contains the *Ti* (tumour inducing) plasmid. This plasmid is able to transfer genetic material into plant cells and causes crown gall disease. Several regions on the plasmid (identified above) help it to infect plants. The plasmid is just over 200,000 bp long and contains 196 genes. The mapping of its genes has made it of great importance in the creation of transgenic plants.

Plasmids have provided a tool with which to introduce novel genetic material into an organism. A new gene can be spliced into a plasmid and the plasmid inserted into a recipient organism (e.g. a bacterium). The bacteria will then produce the product encoded by the gene. This methodology has enabled the industrial-scale production of valuable gene products, such as human insulin, from genetically engineered organisms.

1. What is a plasmid? \_\_\_\_\_
2. Explain how a plasmid can convey a survival advantage to bacteria under certain conditions: \_\_\_\_\_
3. (a) Why are plasmids (such as the *Ti* plasmid) useful to genetic engineers? \_\_\_\_\_
- (b) Into which region of the *Ti* plasmid would you insert a gene in order for it to be transferred into a host plant cell? \_\_\_\_\_

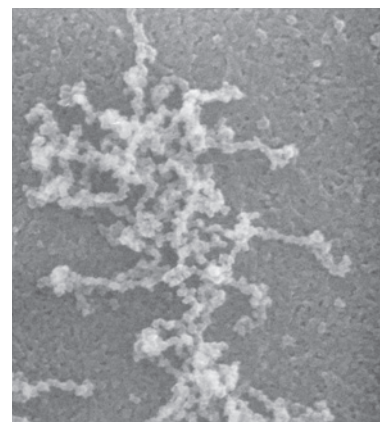
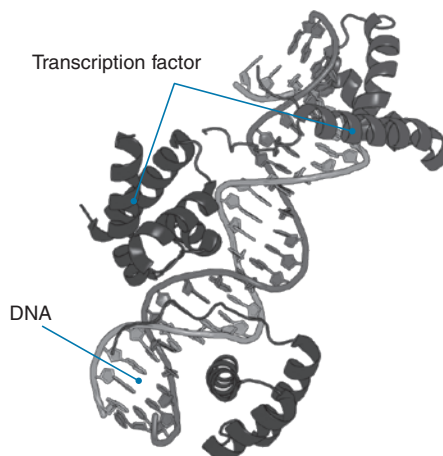


# 167 Controlling Gene Expression

**Key Idea:** Gene expression is tightly regulated. It begins when RNA polymerase attaches to the promoter region of a gene. All the cells in your body contain identical copies of your genetic instructions. Yet these cells appear very different (e.g. muscle, nerve, and epithelial cells have little in common). These morphological differences reflect profound differences

in the expression of genes during the cell's development. For example, muscle cells express the genes for the proteins that make up the contractile elements of the muscle fibre. This diversity of cell structure and function reflects the precise control over the time, location, and extent of expression of a huge variety of genes.

The physical state of the DNA in or near a gene is important in helping to control whether the gene is even available for transcription. When the **heterochromatin** is condensed, the transcription proteins cannot reach the DNA and the gene is not expressed. To be transcribed, a gene must first be unpacked from its condensed state. Once unpacked, control of gene expression involves the interaction of **transcription factors** (shown attached to the DNA molecule right) with DNA sequences that control the specific gene. Initiation of transcription is the most important and universally used control point in gene expression.

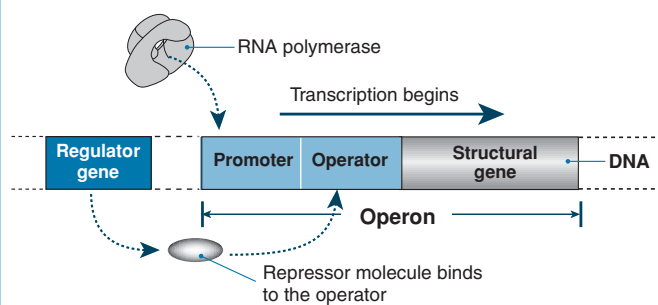


Unravelled mRNA after transcription.

Before transcription can take place, RNA polymerase must attach to the **promoter** region of the DNA. The promoter is upstream of the DNA to be transcribed. The process by which RNA polymerase binds to the promoter is different in prokaryotes and eukaryotes (below).

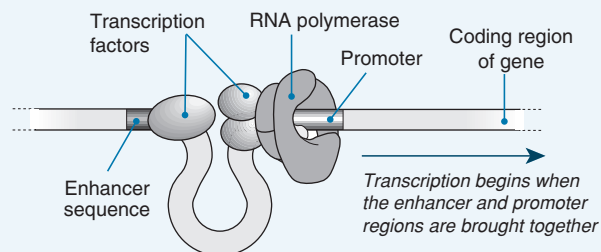
## Control of Gene Expression in Prokaryotes

In **prokaryotes**, the RNA polymerase and associated proteins bind directly to the promoter region. The promoter is normally very close to the DNA region to be transcribed. Transcription of the structural gene is controlled by a **regulator** gene, which produces a repressor molecule that may bind to the operator and block transcription. The promoter, operator, and DNA to be transcribed are called an **operon**.



## Control of Gene Expression in Eukaryotes

Eukaryote genes do not exist as operons. In **eukaryotes**, several **transcription factors** are required to bind the RNA polymerase to the promoter (the RNA polymerase cannot bind directly). Some of these transcription factors may also bind to the **enhancer** region which may be quite distant from the promoter. The transcription factors cause the promoter and enhancer to come together, which initiates transcription.



1. (a) How is transcription initiated in prokaryotes? \_\_\_\_\_

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(b) How is transcription initiated in eukaryotes? \_\_\_\_\_

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2. What is the role of the promoter? \_\_\_\_\_

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# 204 Mechanism of Natural Selection

**Key Idea:** Natural selection is the evolutionary mechanism by which organisms that are better adapted to their environment survive to produce a greater number of offspring.

Evolution is the change in inherited characteristics in a population over generations. Evolution is the consequence

of interaction between four factors: (1) The potential for populations to increase in numbers, (2) Genetic variation as a result of mutation and sexual reproduction, (3) competition for resources, and (4) proliferation of individuals with better survival and reproduction.

**Natural selection** is the term for the mechanism by which better adapted organisms survive to produce a greater number of viable offspring. This has the effect of increasing their proportion in the population so that they become more common. This is the basis of Darwin's theory of evolution by natural selection.

We can demonstrate the basic principles of evolution using the analogy of a 'population' of M&M's candy.



In a bag of M&M's, there are many colours, which represents the variation in a population. As you and a friend eat through the bag of candy, you both leave the blue ones, which you both dislike, and return them to bag.



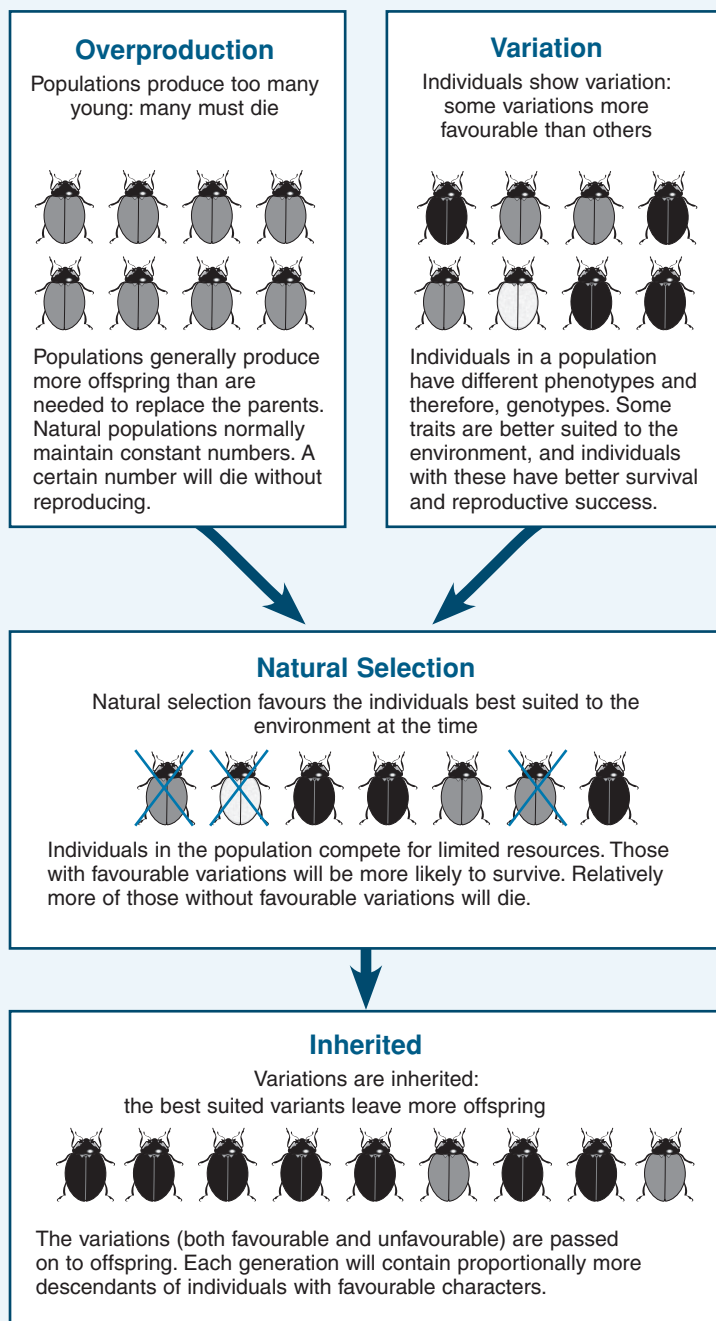
The blue candy becomes more common...



Eventually, you are left with a bag of blue M&M's. Your selective preference for the other colours changed the make-up of the M&M's population. This is the basic principle of selection that drives evolution in natural populations.

## Darwin's Theory of Evolution by Natural Selection

Darwin's theory of evolution by natural selection is outlined below. It is widely accepted by the scientific community today and is one of founding principles of modern science.



1. Identify the four factors that interact to bring about evolution in populations: \_\_\_\_\_

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# 259 Trends in Dentition

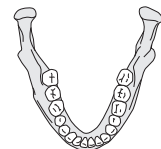
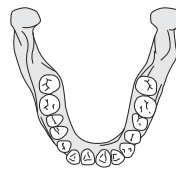
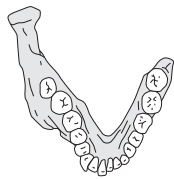
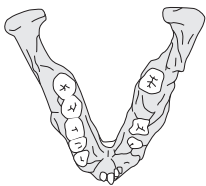
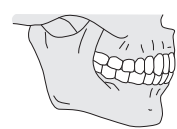
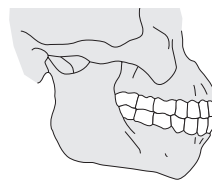
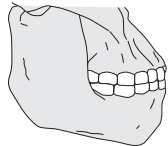
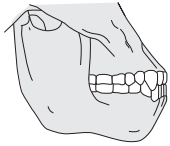
**Key Idea:** Changes in dentition (the type, number, and arrangement of teeth) and jaw structure of our hominin ancestors can reveal information about our evolution.

During early hominin evolution teeth (especially the molars) and jaws tended to be large. The paranthropines are the extreme example of this trend. Their diet of coarse vegetation required very large and powerful jaws and molars. During

the course of hominin evolution, there was a general trend for a reduction in the size of the teeth and jaw. This was a likely consequence of including a greater proportion of cooked foods, which required less chewing, in the diet. The teeth of modern humans are relatively small and generalised, reflecting an omnivorous diet of mainly processed (e.g. cooked) foods.

## Early Hominins

## Late Hominins



### *Australopithecus afarensis*

- Relatively large canine teeth
- Relatively large jaw
- V-shaped dental arcade
- Thin tooth enamel
- Diet probably consisted of fruits with some tougher material

### *Australopithecus africanus*

- Reduced canine teeth
- Large molars
- Dental arcade intermediate between *A. afarensis* and *H. sapiens*
- Thick tooth enamel
- Diet probably included vegetable matter, nuts, seeds, insects, and eggs

### *Homo erectus*

- Thick jaw bones
- No chin
- Relatively large molars
- Parabolic dental arcade
- Thick tooth enamel
- Diet probably included vegetable material and a large proportion of meat

### *Homo sapiens*

- Shortened jaw, allows large bite force to be generated with little effort
- Chin reinforces jaw, but leaves room for tongue muscles
- Thick tooth enamel
- Small molars adapted to chewing cooked and soft food
- Parabolic dental arcade

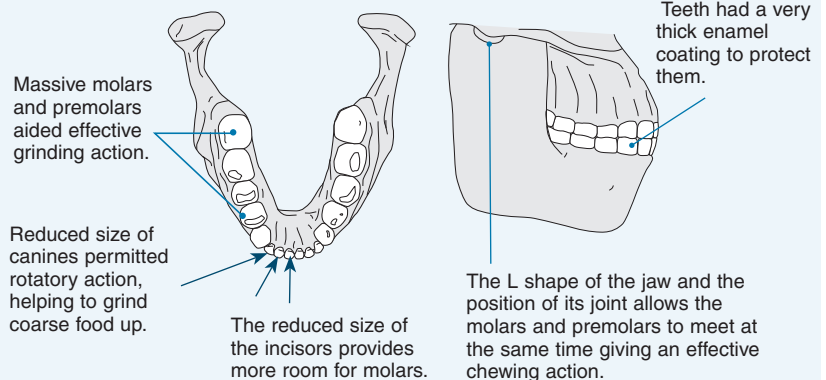
Dental formulae all follow: I-2, C-1, P-2, M-3



In many primates, the canine teeth are used in behavioural and social interactions, especially in species which show marked sexual dimorphism. Threat gestures, such as yawning (above), help maintain social order.

## Adaptations to a Coarse Diet

*Paranthropus boisei* had jaws and teeth adapted to a diet of coarse vegetation and hard seeds. Their jaws produced a massive bite force of 2161 Newtons, which helped to break food up.



1. Describe the general trend in the evolution of hominin teeth: \_\_\_\_\_

2. What is one possible purpose of the chin in modern humans? \_\_\_\_\_

PAGE FEATURES

Activity number

Activities are numbered to make navigation through the workbook easier.

Key idea

Each activity has a key idea summarising its primary focus. The key idea helps students to understand the message of the page.

Activity codes

Page codes indicate the nature of the activity (e.g. KNOW indicates knowledge or basic understanding). Other codes indicate a focus on practical skills, data handling, vocabulary, or comprehension. Activities for reference or assessment are also identified.

Web codes

Weblinks refer to an external URL supporting the material in this activity.

Link codes

Make it easy for students to locate related activities elsewhere in the workbook.

Visual impact

Large annotated diagrams explain the main idea of the activity, supporting and reinforcing the text based content.

Easy to understand content

The content has been written in language that students can easily understand. The content is presented in logically organised blocks, making it easier for students to access and engage with the information.

Direct questioning style

The questions are designed to help students consolidate their understanding of biological processes, principles, and vocabulary. The question style is direct, making the message clear.

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Mutations and Development

**Key Idea:** Errors in DNA replication during meiosis can produce new alleles that can be inherited. Every time DNA is replicated, there is a chance that a copying error will occur. DNA replication has a low **error rate**, with only one mistake for every billion base pairs copied. Changes to the DNA base sequence, called **mutations**, with no effect on the coded product are called **silent**. However, some mutations can create new alleles. Most new alleles will be detrimental. One example is NSRD, a common form of genetic hearing loss, which accounts for up to 50% of childhood deafness.

Chromosome 13

Six connexin 26 peptide chains join together to form a connection between cells. If these proteins are not made correctly, they are unable to form the final structure.

Connexin 26 peptide chains

Normal DNA sequence

Leu Gly Gly Val Asn

28 42

c t t g g g g c t a t a a c

A three base sequence coding for an amino acid is called a **codon**.

Mutated sequence

Leu Gly Val Stop

28 42

c t t g g g g t t g a a c a

TGA codes for stop

The most common mutation in this gene is **deletion of the 35th base (G)**. DNA is read in groups of three bases, so deleting the guanine alters the reading sequence. The result is a short peptide chain, which cannot function correctly and results in deafness.

The deletion of this one base causes the formation of a **recessive allele**. Deafness results when there are two recessive alleles.

**Harmful Mutations**

Most mutations cause harmful effects, usually because they stop or alter the production of a protein (often an enzyme). Albinism (above) is one of the more common mutations in nature, and leaves an animal with no pigmentation.

Arg

G C T

c G A

Normal sequence

G C C

c G G

Mutated sequence

Silent Mutations

Silent mutations do not change the amino acid sequence nor the final protein. In the genetic code, several codons may code for the same amino acid. Silent mutations are also **neutral** if they do not alter the fitness of the organism.

**Beneficial Mutations**

Sometimes mutations help the survival of an organism. In viruses (e.g. *Influenzavirus* above) genes coding for the glycoprotein coat are constantly mutating, producing new strains that avoid detection by the host's immune system.

1. How can changes in a DNA sequence occur? \_\_\_\_\_

2. How can a mutation in a single base be as damaging as a mutation in a sequence of bases? \_\_\_\_\_

3. Explain how mutation can be harmful or beneficial: \_\_\_\_\_

KNOW

166

121

172

176

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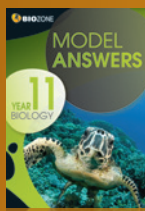
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