

# EBIOLOGY<br/>UNITS 1&2



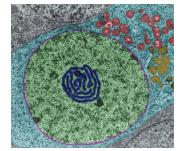


# **Cellular Structure and Function**

	Cells are the unit of life	Activity
	Key skills and knowledge	number
	1 Recognise cells as the basic unit of life on Earth. Describe the characteristics of living organisms and explain why viruses do not fulfil the criteria for being living cells.	9
	2 List the basic biochemical components of cells. Appreciate the role of water in life on Earth and summarise its biologically important properties.	10
	<sup>3</sup> Describe the requirements of cells in terms of their immediate environment. Explain how unicellular and multicellular organisms meet the challenges for surviving in different environments.	11
	4 Describe the main differences between eukaryotic and prokaryotic cells. Recognise the cells of fungi, plants, protists, animals, and bacteria by their characteristic features.	9 12 14
t	5 Use drawings and electron micrographs to compare and contrast the structure of prokaryotic cells and eukaryotic cells.	13 14







#### Limitations to cell size: surface area to volume ratios

	Key skills and knowledge	
	6 Describe the range of cell sizes. Express cell sizes in different units of measurement.	
	7 Describe how cells exchange substances by diffusion. Identify the factors affecting rates of diffusion, explain their effect, and relate these to biological systems.	
	8 <b>PRAC</b> Investigate diffusion across membranes using a model system.	
	9 Explain the importance of surface area to volume ratio in limiting cell size and describe the role of cellular organelles in creating cellular compartments with specific functions.	
	10 <b>PRAC</b> Investigate the effect of cell size on the rate and efficiency of diffusion.	
	11 Explain how cells overcome the limitations to cell size by changes in shape and by the way they are organised in tissues.	
	The structure and specialisation of cell organelles	
	Key skills and knowledge	
	12 Compare and contrast the ultrastructure of plant cells and animal cells in terms of their organelles. Identify these organelles in drawings and in light and electron micrographs.	20-
	13 Describe the specialisations of plant and animal cellular organelles for specific functions.	20-
	The structure and function of the plasma membrane	
	Key skills and knowledge	-
	14 Describe the structure of the plasma membrane and its role as a partially permeable boundary between the internal and external environments of the cell. Recognise that internal membranes, e.g. of membranous organelles, have the same basic structure.	23-
atio	15 PRAC Investigate factors affecting membrane structure and permeability. How might your findings be relevant to survival in different environments?	
	16 Explain the role of the plasma membrane in the movement of substances by diffusion, facilitated diffusion, and active transport (including ion pumps, cotransport, and cytosis). Include an explanation of the movement of water by osmosis. Explain the effects that solutions of different solute concentration can have on plant and animal cells.	16 29-
	17 <b>PRAC</b> Investigate the effects of solutions of different solute concentration on plant cells. Use your results to estimate the osmolarity of a cell, e.g. a potato cell.	

#### Key terms

active transport
carrier protein
cell wall
centrioles
channel protein
chloroplast
cilia
concentration gradient
cytoplasm
diffusion
electron micrograph
endoplasmic reticulum
eukaryotic cell
facilitated diffusion
flagella
Golgi apparatus
hypertonic
hypotonic
ion pump
isotonic
light microscope
lysosome
magnification
mitochondrion
nucleic acid
nucleolus
nucleus
organelle
osmolarity
osmosis
partially permeable
passive transport
phospholipid
plasma membrane
plasmolysis
prokaryotic cell
protein
resolution
ribosome
rough ER (rER)
smooth ER (sER)
surface area: volume ra

turgor

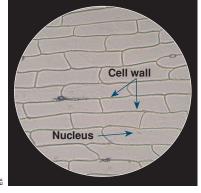
vacuole

## 13 Interpreting Images of Cells

## **Key Idea**: Different microscopy techniques produce different views of cells and their features.

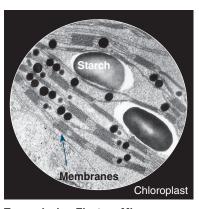
The microscope is an important tool in biology for viewing cells and their features, which are far too small to be seen by the human eye. High power compound light microscopes use visible light and a combination of lenses to magnify objects up to several 100 times. Electron microscopes use beams of

**Dissecting** microscopes are used for dissections, observing microbial cultures, and for identifying and sorting organisms, like this small crustacean.

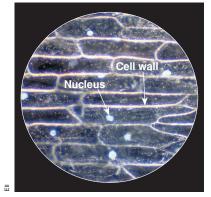


types of atoms.

These onion epidermal cells are viewed with standard **bright field** lighting. Very little detail can be seen. The cell nuclei are barely visible.



**Transmission Electron Microscopes** (TEMs) produce extremely high resolution images of the interior of cells and transparent objects.



electrons and computer imaging to capture extremely fine

detail of either surface or internal cellular features. They can

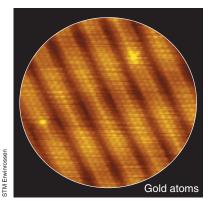
magnify images up to 500,000 times. Scanning Tunnelling

Microscopes (STMs) can magnify object ten times more than

that. With a resolution of 0.1 nanometers, STMs operate at

the edge of the quantum realm and are able to image some

**Dark field** illumination is excellent for viewing specimens that are almost transparent. The nuclei of these onion epidermal cells are clearly visible.



Scanning Tunnelling Microscopes (STMs) produce images based on current variation between an extremely fine needle and the object it moves over.

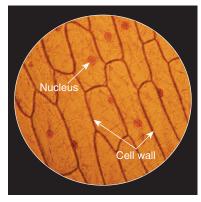


g

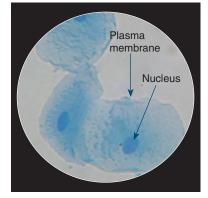
Scanning Electron Microscopes (SEMs) produce extremely high resolution images of the surface of cells and objects.



Some parts of the cell take up stains (chemical dyes) better than others. Stains can be used to highlight parts of the cell for better viewing with a microscope or they can improve contrast. A wide range of chemicals act as stains, including iodine and methylene blue.



lodine is used to increase the contrast in transparent tissues, such as this onion epidermis. Iodine stains are also used to show the presence of starch, binding starch to produce a blue-black colour.



Methylene blue is a positively charged stain commonly used when viewing animal cells. It has a strong affinity for DNA (in the nucleus) and a weaker affinity for RNA (in the cytoplasm).



Some bacteria can be identified and viewed using Gram staining. Bacteria are classed as Gram positive and Gram negative depending on whether or not the stain is retained by the cell wall.

•

#### What is magnification?

Magnification refers to the number of times larger an object appears compared to its actual size.

measured size of the object
actual size of the object
size of the image
magnification

#### What is resolution?

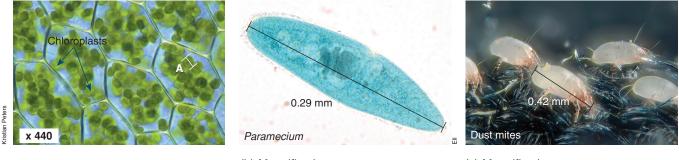
Resolution is the ability to distinguish between close together but separate objects. Resolution is a function of wavelength of light used to view the object. Examples of high and low resolution for separating two objects viewed under the same magnification are given below.

**High resolution** 

Low resolution

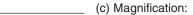


1. Calculate the length or magnification of the object or organism:

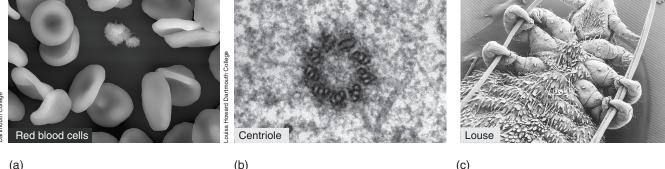


(a) Length of A:

(b) Magnification:



2. Identify which type of microscope (optical microscope, SEM, or TEM) was used to produce each of the images in the photos below (a to f):

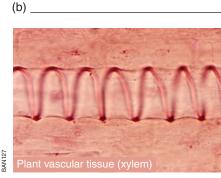


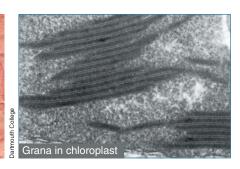
Dartmouth College











(f) (e) \_ 3. Identify the labelled structures: D Blood cells Elodea leaf (c) \_\_\_\_ (b) \_ (d) \_ (a) \_

Dr Graham Be

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## **28** Estimating Osmolarity of Cells

**Key Idea**: Determining loss or gain of mass in tissues allows us to determine the osmolarity of the tissue's cells. The osmolarity (a measure of solute concentration) of a

cell or tissue can be estimated by placing part of the cell or

tissue into a series of solutions of known concentration and observing if the tissue loses (hypertonic solution) or gains (hypotonic solution) water. The solution in which the tissue remains unchanged indicates the osmolarity of the tissue.



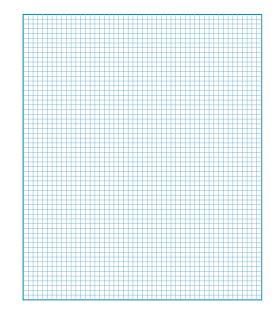
Investigation 2.3 Estimating osmolarity

See appendix for equipment list.

- 1. Prepare 6 beakers of sucrose ( $C_{12}H_{22}O_{11}$ , table sugar) solution with the concentrations of 0.0 (distilled water), 0.1, 0.2, 0.3, 0.4, and 0.5 mol/L of sucrose (0, 34.2 g , 68.5 g, 102.6 g, 136.9 g, and 171.1 g per litre). Label the beakers so that they can be easily identified at the end of the experiment.
- 2. Peel a potato and cut it into 18 identical cubes 1 cm<sup>3</sup> (1 cm x 1 cm x 1 cm) or use a cork borer to produce 18 identical cylinders of potato. Pat the potato cubes dry with a paper towel.
- 3. Weigh three cubes together, record their mass in the table below under initial mass. Place the cubes in the beaker of distilled water.
- 4. Repeat step 3 with the other 15 potato cubes and concentrations. Make sure you identify each beaker so the cubes can be weighed at the end of the experiment.
- 5. Leave the potato cubes in the solutions for at least 40 minutes (or up to 24 hours).
- 6. Remove the potato cubes from the distilled water and pat dry with a paper towel. Weigh all three together and record their mass in the table below under final mass.
- 7. Repeat for all the other concentrations of sucrose.
- 8. Calculate the change in mass (if any) for all the concentrations. Then calculate the % change (+ or -) (this removes any error based on the masses of the potato cubes not being identical).
- 9. Plot the % change vs sucrose concentration on the grid provided.

Sucrose concentration (mol/L)	Initial mass (I) (g)	Final mass (F) (g)
0.00		
Change (C) (F-I) (g)		
% Change (C/I x 100)		
0.1		
Change (C) (F-I) (g)		
% Change (C/I x 100)		
0.2		
Change (C) (F-I) (g)		
% Change (C/I x 100)		
0.3		
Change (C) (F-I) (g)		
% Change (C/I x 100)		
0.4		
Change (C) (F-I) (g)		
% Change (C/I x 100)		
0.5		
Change (C) (F-I) (g)		
% Change (C/I x 100)		

1. Use the grid below to draw a line graph of the sucrose concentration vs total % change in mass:



- 2. Use the graph to estimate the osmolarity of the potato (the point where there is no change in mass):
- 3. Which of the solutions are hypotonic? Which are hypertonic?



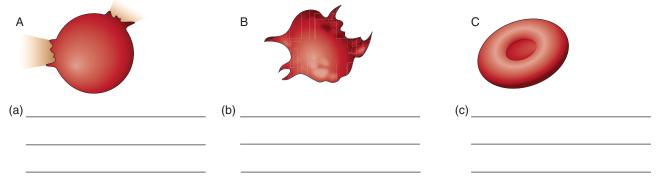
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## 34 Chapter Review: Did You Get It?

#### 1. Match each term to its definition, as identified by its preceding letter code.

active transport	A	The energy-requiring movement of substances across a biological membrane against a concentration gradient.
concentration gradient	в	Movement of substances across a biological membrane without energy expenditure.
diffusion	С	The passive movement of molecules from high to low concentration.
osmosis	D	The gradual difference in the concentration of solutes in a solution between two regions. In biology, this usually results from unequal distribution of ions across a membrane.
passive transport	E	Passive movement of water molecules across a partially permeable membrane down a concentration gradient.

2. The diagrams below depict what happens when a red blood cell is placed into three solutions with differing concentrations of solutes. Describe the tonicity of the solution (in relation to the cell) and describe what is happening:



- 3. Explain how the properties of the phospholipid molecule result in the bilayer structure of membranes:
- 4. Using the formulae: cuboid SA = 2(lh + lw + hw), cuboid volume = lwh, calculate the surface area to volume ratio of the following cell shapes:

- (a) A cubic cell 6 μm x 6 μm x 6 μm: \_\_\_\_
- (b) A cuboid cell 1 μm x 12 μm x 5 μm: \_\_\_\_
- (c) Which of these cells would exchange substances with its environment most efficiently and why: \_\_\_\_\_
- 5. Consider the two diagrams below. For each, draw in the appropriate box what you would expect to see after one hour.

Particle with diameter of 5 nm	Particle with diameter of 20 nm	After one hour:
Soluble particles placed in at high concentration	Container of water at 20° C	
	Partially permeable membrane with pores of 10 nm.	

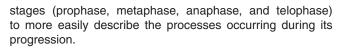
## **39** Recognising Stages in Mitosis

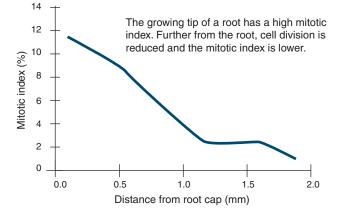
**Key Idea**: The stages of mitosis can be recognised by the organisation of the cell and chromosomes. Although mitosis is a continuous process it is divided into four

The mitotic index

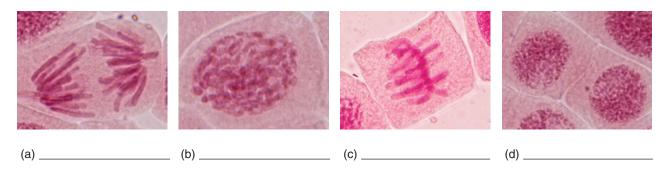
The mitotic index measures the ratio of cells in mitosis to the number of cells counted. It is a measure of cell proliferation and can be used to diagnose cancer (because cancerous cells divide very quickly). In areas of high cell growth the mitotic index is high such as in plant apical meristems or the growing tips of plant roots. The mitotic index can be calculated using the formula below:

Mitotic index = Number of cells in mitosis Total number of cells





1. Use the information in the previous activity to identify which stage of mitosis is shown in each of the photographs below:



2. (a) The light micrograph (right) shows a section of cells in an onion root tip. These cells have a cell cycle of approximately 24 hours. The cells can be seen to be in various stages of the cell cycle. By counting the number of cells in the various stages it is possible to calculate how long the cell spends in each stage of the cycle. Count and record the number of cells in the image that are in mitosis and those that are in interphase. Cells in cytokinesis can be recorded as in interphase. Estimate the amount of time a cell spends in each phase.

Stage	No. of cells	% of total cells	Estimated time in stage
Interphase			
Mitosis			
Total		100	

- (b) Use your counts from 2(a) to calculate the mitotic index for this section of cells.
- 3. What would you expect to happen to the mitotic index of a population of cells that loses the ability to divide as they mature?

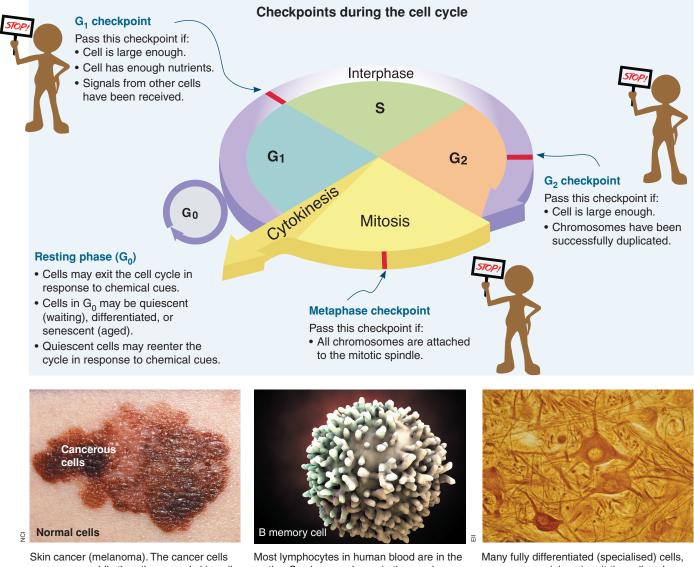
38

Onion root tip cells



# 40 Regulation of the Cell Cycle

**Key Idea**: Regulatory checkpoints are built into the cell cycle to ensure that the cell is ready to proceed from one phase to the next. The failure of these systems can lead to cancer. Cell checkpoints give cells a way to ensure that all cellular processes have been completed correctly before entering the next phase. There are three checkpoints in the cell cycle. At each checkpoint, a set of conditions determines whether or not the cell will continue into the next phase. Cancer can result when the pathways regulating the checkpoints fail. Non-dividing cells enter a resting phase ( $G_0$ ), where they may remain for a few days or up to several years. Under specific conditions, they may re-enter the cell cycle.



grow more rapidly than the normal skin cells because normal cell regulation checkpoints are ignored. This is why the cancerous cells sit higher than the normal cells and can rapidly spread (a process called metastasis). Most lymphocytes in human blood are in the resting  $G_0$  phase and remain there unless they are stimulated by specific antigens to reenter the cell cycle via  $G_1$ .  $G_0$  phase cells are not completely dormant, continuing to carry out essential cell functions in reduced form.

Many fully differentiated (specialised) cells, e.g. neurones (above), exit the cell cycle permanently and stay in  $G_0$ . These cells continue their functional role in the body, but do not proliferate. Senescent cells have accumulated mutations, lose function, and die.

- 1. Explain the importance of cell cycle checkpoints:
- 2. In terms of the cell cycle and the resting phase (G<sub>0</sub>), distinguish between the behavior of fully differentiated cells, such as neurones, and cells that are quiescent, such as B memory cells



## **125** Adaptations of Xerophytes

**Key Idea**: Xerophytes are plants with adaptations that allow them to conserve water and survive in dry environments. Plants adapted to dry conditions are called **xerophytes**. Xerophytes are found in a number of environments, but all

- Most xerophytes are found in deserts, but they may be found in humid environments, provided that their roots are in dry micro-environments (e.g. the roots of epiphytic plants that grow on tree trunks or branches).
- Many xerophytes have a succulent morphology. Their stems are often thickened and retain a large amount of water in the tissues, e.g. *Aloe*.
- Many xerophytes have a low surface area to volume ratio, reducing the amount of water lost through transpiration.
- Salt tolerant plants and many alpine species may show xeromorphic features in response to the lack of free water and high transpirational losses in these often windy, exposed environments.



absence of leaves.

Acacia trees have deep root systems, allowing them to draw water from sources deep underground.

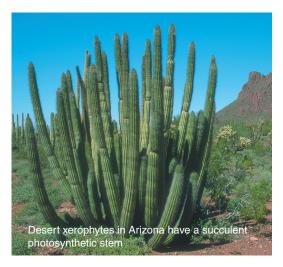


show adaptations to conserve water. These adaptations

include small, hard leaves, an epidermis with a thick cuticle,

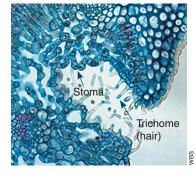
sunken stomata, succulence, and permanent or temporary

An outer surface coated in fine hairs traps air close to the surface and reduces the transpiration rate.





Grasses on coastal sand dunes (e.g. marram grass, above) curl their leaves. Stomata are sunken in pits, creating a moist microclimate around the pore, which reduces transpiration rate.



Oleander has a thick multi-layered epidermis and the stomata are sunken in trichome-filled pits on the leaf underside which restrict water loss. Trichomes (leaf "hairs) maintain a layer of still air at the leaf surface.

What is a xeromorphic adaptation? \_\_\_\_\_

2. Describe three xeromorphic adaptations of plants that reduce water loss:

- 3. (a) How does creating a moist microclimate around the areas of water loss reduce the transpiration rate?

(b) How do trichomes contribute to reducing the transpiration rate?

4. How does a low surface area to volume ratio in a plant such as a cactus reduce water loss? \_\_\_\_

5. How does a cactus photosythesise given it has no leaves? \_\_\_\_\_



#### Adaptations in cacti

Desert plants, such as cacti (below), must cope with low or sporadic rainfall and high transpiration rates.



Leaves modified into spines or hairs to reduce water loss. Light coloured spines reflect solar radiation.

Rounded shape reduces surface area.

Stem becomes the major photosynthetic organ, plus a reservoir for water storage.

The surface tissues of many cacti are tolerant of temperatures over 50°C.

Cacti have a shallow, but extensive fibrous root system. When in the ground the roots are spread out around the plant.

### DID YOU KNOW?

Australia has no native cacti although many have become naturalised. However, two endemic species, the Wongan cactus (*Daviesia euphorbioides*) and the Dunna Dunna (*Lawrencia helmsii*) resemble cacti and are often mistaken for them (an example of convergence, opposite).

#### Convergent adaptation in unrelated xerophytes

- The North American cactus and African Euphorbia species shown above are both xerophytes. They have evolved similar structural adaptations to conserve water and survive in a hot, dry, desert environment. Although they have a similar appearance, they are not related. They provide an excellent illustration of how unrelated organisms living in the same environment have independently evolved the same adaptations to survive.
- Their appearance is so similar at first glance that the Euphorbia is often mistaken for a cactus. Both have thick stems to store water and both have lost the presence of obvious leaves. Instead, they have spines or thorns to conserve water (a leafy plant would quickly exhaust its water reserves because of losses via transpiration). In cacti, spines are highly modified leaves. In Euphorbia, the thorns are modified stalks. It is not until the two flower that their differences are obvious.



6. (a) Explain why the North American cactus and African Euphorbia species have evolved such similar adaptations:

(b) Identify two ways in which a cactus and a euphorbia can be distinguished:

7. Why would a shallow fibrous root system be an advantage to a cactus? \_\_\_\_\_

8. Identify the features of the xerophyte shown that help it conserve water:



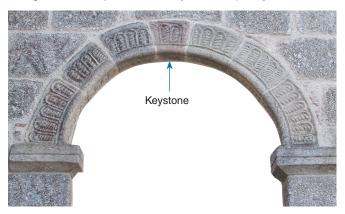
## **133** Interdependencies: Animal Keystone Species

**Key Idea**: All organisms within an ecosystem contribute to its structure and functioning, but keystone species have a disproportionate effect on ecosystem processes.

Although every species has a role in ecosystem function, some have a disproportionate effect on ecosystem processes and stability (how unchanging the ecosystem is over time). These species are called **keystone species** and they are important

#### Why are keystone species important?

A keystone species is one that plays a unique and crucial role in the way an ecosystem functions. Often, but not always, keystone species are top predators. The role of the keystone species varies from ecosystem to ecosystem, but the loss of a keystone species from any ecosystem has a domino effect, and a large number of species can be affected. This can lead to can rapid ecosystem change or the collapse of the ecosystem completely.



The term keystone species comes from the analogy of the keystone in a true arch (above). An archway is supported by a series of stones, the central one being the **keystone**. If the keystone is removed the arch collapses. because they play a pivotal role in the way the ecosystem works, e.g. as top predators or by recycling nutrients. The loss of a keystone species can have a large and rapid impact on the structure and function of an ecosystem, changing the balance of relationships and leading to instability. This has important implications for ecosystem management because many keystone species are endangered.



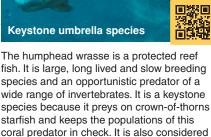
#### Keystone species in action

The idea of the keystone species was first hypothesised in 1969 by Robert Paine. He studied an area of rocky seashore, noting that diversity seemed to be correlated with the number of predators (ochre starfish) present (i.e. diversity declined as the number of predators declined).

To test this he removed the starfish from an 8 m by 2 m area of seashore. Initially, the barnacle population increased rapidly before collapsing and being replaced by mussels and gooseneck barnacles. Eventually the mussels crowded out the gooseneck barnacles and the algae that covered the rocks. Limpets that fed on the algae were lost and the number of species present in the study area dropped from 15 to 8.



The endangered southern cassowary is a keystone species in Australia's wet tropics. They are obligate fruit eaters, and their gut passes seeds, unharmed, into a pile of manure. More than 200 plant species depend on the cassowary to disperse their seeds, yet their populations are all declining. Their loss would also mean the loss of an ecological role.



an umbrella species because its protection

benefits a large number of other species.



Top predators, such as Australia's dingo, are often keystone species. Many conservationists regard dingoes as a functional replacement for native predators that are now extinct, such as the Tasmanian tiger. Dingoes have a varied diet and are a major constraint on introduced species, such as a foxes and pigs, thereby helping to maintain native mammal diversity.

#### 1. Why are keystone species so important to ecosystem function? \_





Many sharks are top predators and are keystone species in the waters around Australia. One shark species inhabiting Shark Bay (WA) is the tiger shark. It doesn't even have to kill its prey to exert an effect on ecosystem structure. The presence of the tiger shark causes marine herbivores such as green turtles and dugongs to avoid the area or to spend less time grazing because they are looking out for the sharks. As a result, the seagrass meadows thrive and support many more species than would be possible if they were grazed intensively by herbivores. As a result, biodiversity in Shark Bay is high. Fishing is the main threat to tiger sharks as they hunted for their flesh, fins, and skin. Finning is banned in Australian waters, but still continues illegally because of the difficulty in policing.



The grey-headed flying fox (*Pteropus poliocephalus*) is found in a variety of habitats along the east coast of Australia, including Victoria. The grey-headed flying fox feeds on the fruit and nectar of over 180 species of trees, including Australian natives *Eucalyptus*, *Banksia*, palms, and myrtles. It will fly up to 50 km each night looking for food and this allows it to fulfill an important ecological role by dispersing the pollen and seeds of a wide range of plants. Its role is especially important in the subtropical rainforests as it is the only mammalian species to consume nectar and fruit in these regions. The species is under threat from the loss of foraging and roosting habitat and control measures by horticulturists to prevent crop losses.

2. For each species below, summarise the features of its ecology that contribute to its position as a keystone species:

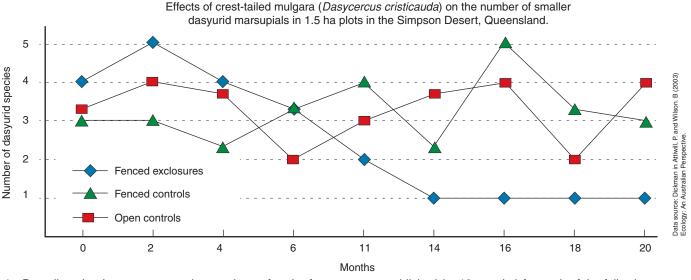
(a)	Southern cassowary:
(b)	Humphead wrasse:
(c)	Dingo:
(പ)	
(u)	Tiger shark:
(a)	Crow boarded flying for:
(e)	Grey-headed flying fox:

3. The arrival of the dingo in Australia probably resulted in the loss or reduction of many top marsupial predators. It now plays a role as a keystone species in modern Australia. How might this have occurred?

#### Case study: crest-tailed mulgara as a keystone species

- Two species of mulgara (*Dasycercus* genus) are found in Australia, the brush-tailed mulgara and the crest-tailed mulgara. Mulgara are nocturnal marsupials belonging to the family Dasyuridae, which includes the Tasmanian devil and the quolls. Both mulgara species are small (30 cm long from head to tail) and weigh up to 190 g.
- Mulgaras live in arid central Australia, and burrow 50 cm under the surface to avoid the heat. While the brush-tailed mulgara has an extensive range through the middle of Australia, the crest-tailed mulgara is found only in a small part of the Simpson Desert within Queensland's borders.
- The effect of the crest-tailed mulgara (*Dasycercus cristicauda*) as a keystone species was tested by excluding them from a 1.5 ha plot of land. Fenced exclosures were established 10 months after sampling began. All dasyurid species (except mulgara) could access the site. Fenced controls and open controls were established at the same time. All dasyurid species (including mulgara) could access these plots. The results are shown in the graph below.





- 4. Describe what happens to species numbers after the fences were established (at 10 months) for each of the following:
  - (a) Fenced exclosures:\_
  - (b) Fenced controls: \_\_\_\_
  - (c) Open controls: \_\_\_\_\_
- 5. Describe the difference in species numbers between the fenced exclosure and the:
  - (a) Fenced control: \_\_\_\_\_
  - (b) Open control: \_\_\_\_\_
- 6. (a) Based on the data presented above, do you think the crest-tailed mulgara acts as a keystone species?\_\_\_\_
  - (b) Explain your answer: \_\_\_\_

7. Why do you think the researchers included a fenced control and open control?

8. Why let other dasyurid species into the exclosures?\_\_\_\_\_

## Identifying Key Science Skills and Online Support

The tab system is a useful way to identify important parts of the **VCE Biology study design (2022 - 2026)**. The colour coded page tabs show where "**Key Science Skills**" are addressed. The tabs also allow you to see at a glance if online support is provided and if there are content or concept links with other activities.

#### **KEY SCIENCE SKILLS**



## Develop aims and questions, formulate hypotheses and make predictions

- Identify, research, and construct aims and questions
- Identify variables in controlled experiments
- Formulate hypotheses to focus investigation
- Predict possible outcomes



#### Plan and conduct investigations

- Determine methodology, classification and identification, controlled experiments, correlational studies, fieldwork, literature review, modelling, simulations
- Design and conduct investigations, considering procedures, error, amount of data
- Work independently and collaboratively adapt methodology



## Comply with safety and ethical guidelines

- Demonstrate safe lab practices using risk
  assessments
- Apply relevant health and safety guidelines
- Demonstrate ethical conduct when undertaking and reporting investigations



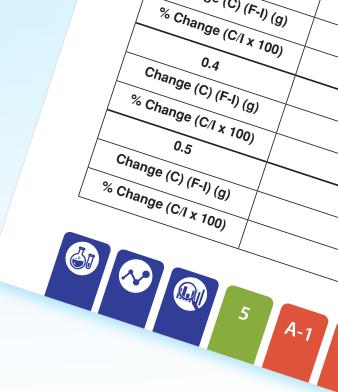
#### Generate, collate, and record data

- Systematically generate and record primary data, collate secondary data
- Record and summarise data, including use of a logbook
- Organise and present data
- Plot graphs showing liner and non-linear relationships



## Analyse and evaluate data and investigation methods

- Process quantitative data using mathematics
- Identify and analyse experimental data qualitatively
- · Identify outliers and contradictory data
- Repeat experiments and evaluate methods





## Construct evidence-based arguments and draw conclusions

- Distinguish scientific and non-scientific ideas
- Evaluate data for evidence-based support of aims, prediction, or hypothesis
- Use reasoning to construct scientific arguments and to draw and justify conclusions
- Identify and describe limitations of conclusions
- Discuss implications of findings



## Analyse, evaluate and communicate scientific ideas

- Use appropriate terminology
- Discuss relevant biological information, ideas, concepts, theories, and models
- Analyse and explain how models and theories are used
- Critically evaluate and interpret scientific and media texts
- Analyse and evaluate bioethical issues
- Use clear and concise expression
- Acknowledge sources of information

#### **ONLINE SUPPORT**

**Resource Hub** 



This tab indicates the activity is supported with online content, through the Resource Hub.

Easy navigation to video clips, animations, 3D models, databases, and spreadsheets to support the activities in the book.

## www.**BIOZONE**.com/au/VCE



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