The Friary Sixth Form



Biology Summer Project Pack 2023



Summer Tasks



Complete the examination questions on the following pages which will help you consolidate the work covered in Year 12.

There are keywords that <u>must</u> be included in the responses in order for marks to be awarded. These are given to you in the space for your responses.

Q1.

Helicase	
Template	
RNA polymerase	
Splicing	
) Describe the structure of proteins	
) Describe the structure of proteins.	
Polymer	
Amino Acid	
Amino Acid	

				(Total 15 ma
(a)	In humans, the e	zyme maltase breaks dow	vn maltose to glucose.	
	This takes place	t normal body temperatur	vn maltose to glucose. e.	
	This takes place Explain why male only breaks	t normal body temperaturo se: down maltose	e.	
	This takes place Explain why male only breaks	t normal body temperaturo se:	e.	
	This takes place Explain why male only breaks	t normal body temperaturo se: down maltose	e.	
	This takes place Explain why mal only break allows this	t normal body temperaturo se: down maltose	e.	
	This takes place Explain why male only breake allows this Tertiary structure	t normal body temperaturo se: down maltose	e.	
	This takes place Explain why malton only breaks allows this Tertiary structure Complementary Catalyst	t normal body temperaturo se: down maltose	e.	
	This takes place Explain why male only breake allows this Tertiary structure Complementary	t normal body temperaturo se: down maltose	e.	
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	This takes place Explain why malton only breaks allows this Tertiary structure Complementary Catalyst	t normal body temperaturo se: down maltose	e.	
	This takes place Explain why malton only breaks allows this Tertiary structure Complementary Catalyst	t normal body temperaturo se: down maltose	e.	

(c)

Describe how proteins are digested in the human gut.

Drawart C C		-
Prevent E-S		_
Shape		
Allosteric site		-
[substrate]		-
		-
		=
		-
		-
		_
		=
	(Tot	al 10 ma
	tructures of starch and cellulose molecules are related to their	
(a) Describe how the s		-
(a) Describe how the structions.		-
(a) Describe how the structions. Shape Osmosis		-
(a) Describe how the structions. Shape Osmosis Size		-
(a) Describe how the structions. Shape Osmosis		-
(a) Describe how the structions. Shape Osmosis Size		-
(a) Describe how the structions. Shape Osmosis Size		-
(a) Describe how the structions. Shape Osmosis Size		-
(a) Describe how the structions. Shape Osmosis Size		-
(a) Describe how the structions. Shape Osmosis Size		-
(a) Describe how the structions. Shape Osmosis Size		-

Scientists have investigated the effects of competitive and non-competitive inhibitors of the enzyme maltase.

Describe the processes involved in the transport of sugars in plant stems.

Q5.

icelles						_
le salts						
atty acids						
ffusion						
						_
						_
Describe how	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	_
	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	_
Describe how	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	_
	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	_
Primary	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	_
Primary Sequence	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	
Primary Sequence Secondary	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	——————————————————————————————————————
Primary Sequence Secondary Bonding	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	
Primary Sequence Secondary Bonding	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	
Primary Sequence Secondary Bonding	the structure of	a protein depe	nds on the am	nino acids it c	ontains.	

(5)

Q6.

(a) Explain **five** properties that make water important for organisms.

Describe the biochemical tests you would use to confirm the presence of lipid, non-reducing sugar and amylase in a sample. Ethanol Benedict's Acid	1etabolite					
Describe the biochemical tests you would use to confirm the presence of lipid, non-reducing sugar and amylase in a sample. Ethanol Benedict's Acid	olvent					
Describe the biochemical tests you would use to confirm the presence of lipid, non-reducing sugar and amylase in a sample. Ethanol Benedict's Acid	leat capacity					
Describe the biochemical tests you would use to confirm the presence of lipid, non-reducing sugar and amylase in a sample. Ethanol Benedict's Acid	atent heat of					
reducing sugar and amylase in a sample. Ethanol	vaporisation					
Ethanol Benedict's Acid						
Ethanol Benedict's Acid						
Ethanol Benedict's Acid						
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Ethanol Benedict's Acid						
Ethanol Benedict's Acid						
Ethanol Benedict's Acid						
Ethanol Benedict's Acid						
Ethanol Benedict's Acid	Describe the	niochemical tests vou w	ould use to confirm	the presence of	f linid non-	
Benedict's Acid	Describe the reducing suga	piochemical tests you w ar and amylase in a sam	ould use to confirm	n the presence o	f lipid, non-	
Acid	reducing suga	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
·	reducing suga	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
Heat	reducing suga	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
	reducing suga Ethanol Benedict's	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
	reducing suga Ethanol Benedict's Acid	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
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	reducing suga Ethanol Benedict's Acid	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
	reducing suga Ethanol Benedict's Acid	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
	reducing suga Ethanol Benedict's Acid	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	
	reducing suga Ethanol Benedict's Acid	piochemical tests you war and amylase in a sam	iple.		f lipid, non-	

Condensation				
Hydrolysis				
Bonds				
	<i>)</i>			
				(Total 45
				(Total 15
mitochondria fr	xplain how cell fractior om a suspension of an	nation and ultracent imal cells.	rifugation can be use	
Describe and emitochondria fromogenisation	xplain how cell fractior om a suspension of an	nation and ultracent imal cells.	rifugation can be use	
mitochondria fr	xplain how cell fractior om a suspension of an	nation and ultracent imal cells.	rifugation can be use	
mitochondria fr	xplain how cell fractior om a suspension of an	nation and ultracent imal cells.	rifugation can be use	
mitochondria fr Homogenisation Filter Isotonic	xplain how cell fractior om a suspension of an	nation and ultracent imal cells.	rifugation can be use	
mitochondria fr Homogenisation Filter Isotonic Cold	xplain how cell fractior om a suspension of an	nation and ultracent imal cells.	rifugation can be use	
mitochondria fr Homogenisation Filter Isotonic	xplain how cell fractior om a suspension of an	ation and ultracent imal cells.	rifugation can be use	
mitochondria fr Homogenisation Filter Isotonic Cold	xplain how cell fractior om a suspension of an	ation and ultracent imal cells.	rifugation can be use	

Describe the chemical reactions involved in the conversion of polymers to monomers and monomers to polymers.

Electrons					
Wavelength					
Resolution					
vacuum					
- decoin	<u> </u>				
					(Total 10
					(Total 10
					(Total 10
Describe how m	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds Template	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase Phosphodiester	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase	nRNA is form	ed by transcr	iption in eul	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase Phosphodiester	nRNA is form	ed by transcr	iption in euk	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase Phosphodiester	nRNA is form	ed by transcr	iption in euk	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase Phosphodiester	nRNA is form	ed by transcr	iption in euk	karyotes.	(Total 10
Hydrogen bonds Template Nucleotides Polymerase Phosphodiester	nRNA is form	ed by transcr	iption in euk	karyotes.	(Total 10

Describe the principles and the limitations of using a transmission electron microscope to investigate cell structure.

Ribosome				
Codon				
Anticodon				
tRNA				
amino acid				
Define 'gene r	nutation' and explain	how a gene mut:	ation can have:	
		now a gene mate	ation can nave.	
	on an individual e effect on an individu	ual.		
Danamarata				
Degenerate				
Tertiary structur				
Increased surviva	ı)			
				(Total 15 ma

Describe how a polypeptide is formed by translation of mRNA.

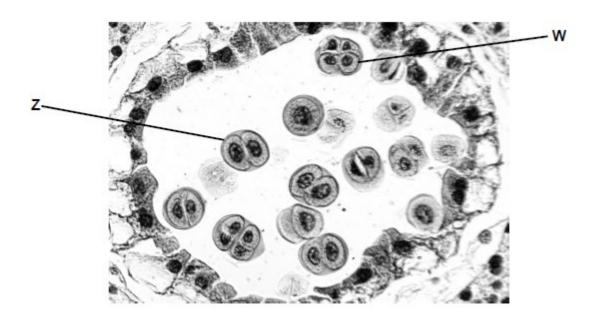
Q9.

(a) Contrast how an optical microscope and a transmission electron microscope work **and** contrast the limitations of their use when studying cells.

Electrons/ light	—		
Resolution			
Dead / alive			
Colour			
	_		

(b) The diagram shows an image from an optical microscope of meiosis occurring in a flower bud of a flowering plant. **W** and **Z** are undergoing meiosis.

(6)



Meiosis	
Haploid / diploid	
,	
A	
	ntal scientist investigated a possible relationship between air pollution and ds produced by one species of tree.
	ed with a very large number of seeds collected from a population of trees in city and also a very large number of seeds collected from a population of untryside.
	ne should collect and process data from these seeds to investigate whether
Describe how	rence in seed size between these two populations of trees.
Describe how	
Describe how there is a differ	
Describe how there is a difference Random	
Describe how there is a different Random Large sample	

(5) (Total 15 marks)

Q10.

Describe	the app							
Prophase								
Metaphase	9							
Anaphase								
Telophase								
Describe variation.	and exp	ain the pr	rocesses t	hat occur c	uring meio	sis that incl	rease ger	netic
Describe variation. Independer segregation	nt	lain the pr	rocesses t	hat occur c	uring meio	sis that inc	rease ger	netic
variation.	nt n	lain the pr	rocesses t	hat occur c	uring meio	sis that inc	rease ger	netic
Independer segregation	nt n	lain the pr	rocesses t	hat occur c	uring meio	sis that inc	rease ger	netic
Independer segregation	nt n	lain the pr	rocesses t	hat occur c	uring meio	sis that inci	rease ger	netic
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Independer segregation	nt n	lain the pr	rocesses t	hat occur o	uring meio	sis that inci	rease ger	netic
Independer segregation	nt n	lain the pr	rocesses t	hat occur o	luring meio	sis that inci	rease ger	netic
Independer segregation	nt n	lain the pr	rocesses t	hat occur o	luring meio	sis that inci	rease ger	netic

Q11.

Semi-	}	
conservative		
Base pairing		
Template		
)	
Describe the beha production of two	aviour of chromosomes during mitosis and explain how this results i genetically identical cells.	in the
Identical		
Identical Chromatids		
Chromatids		
Chromatids Middle		
Chromatids Middle Spindle fibres		

(c)	A cancerous tumour is formed by uncontrolled mitotic division. This results in a mass cells with an inadequate blood supply. Drugs are being developed which only kill cells low oxygen environment. Suggest how these drugs could be useful in the treatment or cancer.	in a
	(Tota	(2 I 13 marks
Q12.		
Diffe	erent cells in the body have different functions.	
(a)	Some white blood cells are phagocytic. Describe how these phagocytic white blood codestroy bacteria.	ells
	Antigens	
	Engulf	
,	Vesicle	
	Lysosome	
	hydrolysis	
		(4
(b)	The epithelial cells that line the small intestine are adapted for the absorption of gluco	
	Explain how.	
(Microvilli	
	Mitochondria	
	Carrier proteins	
	Channel proteins	

Q13.

Bacterial meningitis is a potentially fatal disease affecting the membranes around the brain. *Neisseria meningitidis* (Nm) is a leading cause of bacterial meningitis.

(a)	In the UK, children are vaccinated against this disease. Describe how vaccination can lead to protection against bacterial meningitis.	
	Antigen binds T cell	
	Cytokine	
	B cells	
	Antibodies	
	Memory cell	
		(6)
(b)	Penicillin has been the antibiotic of choice for the treatment of bacterial meningitis. Since the year 2000, strains of <i>Neisseria meningitidis</i> that are resistant to penicillin, sulfonamides and rifampin have been discovered in the UK.	
	Describe how a population of <i>Neisseria meningitidis</i> (Nm) can become resistant to the antibiotics.	se
	Mutation	
	Allele	

	Size		
	Genetic material		
	Ribosomes		
	mitochondria		
			(5)
			(Total 15 marks)
Q14.			
(a)	When a vaccine is g	ven to a person, it leads to the production of antiboo	lies against a
	disease-causing org	nism. Describe how.	
	T cell		
	<u> </u>		
	Cytokine		
	B cell		
	antibody		
'			
			(5)

Contrast the structure of a bacterial cell and the structure of a human cell.

(c)

1	Memory cells										
	Plasma cells										
	Time interval										
	longevity										
`											
										(i otai	10 mark
Q15. (a)	Glucose is abso	rbed from	the lu	men of [.] lium ion	the sm s is inv	all intes	tine into e the abso	pithelial	cells.	e bv	
	Explain how the epithelial cells. Active transport	rbed from transport	the lu	men of lium ion	the sm s is inv	all intes olved in	tine into e the absoi	pithelial rption of	cells. f glucose	e by	
	Explain how the epithelial cells. Active transport Na+	rbed from transport	the lu	men of lium ion	the sm s is inv	all intes olved in	tine into e the absor	pithelial rption of	cells. f glucose	e by	
	Explain how the epithelial cells. Active transport	rbed from transport	the lu	men of lium ion	the sm s is inv	all intes olved in	tine into e the absor	pithelial rption of	cells. f glucose	e by	
	Explain how the epithelial cells. Active transport Na+	rbed from transport	the lu	men of	the sm s is inv	all intes olved in	tine into e the absor	pithelial rption of	cells. f glucose	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+]	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells. f glucose	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+] Facilitated	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells. f glucose	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+] Facilitated diffusion	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells.	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+] Facilitated diffusion	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells.	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+] Facilitated diffusion	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells.	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+] Facilitated diffusion	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells.	e by	
	Explain how the epithelial cells. Active transport Na+ Lower [Na+] Facilitated diffusion	rbed from transport	the lu	men of	the sm s is inv	all intes	tine into e the absor	pithelial rption of	cells.	e by	

Describe the difference between active and passive immunity.

Solubility		
Polarity		
Size		
)	
		(Total 1
		(Total 1
Many different Describe how	substances enter and leave a cell by crossing substances can cross a cell surface membrand	its cell surface membrane
Many different Describe how	substances enter and leave a cell by crossing substances can cross a cell surface membrand	its cell surface membrane
Many different Describe how	substances enter and leave a cell by crossing substances can cross a cell surface membrand	its cell surface membrane
Describe how	substances enter and leave a cell by crossing substances can cross a cell surface membrand	its cell surface membrane
Describe how a Diffusion Lipid-soluble	substances enter and leave a cell by crossing substances can cross a cell surface membrane	its cell surface membrane
Describe how and Diffusion Lipid-soluble Osmosis	substances enter and leave a cell by crossing substances can cross a cell surface membrane	its cell surface membrane
Describe how a Diffusion Lipid-soluble	substances enter and leave a cell by crossing substances can cross a cell surface membrane	its cell surface membrane
Describe how and Diffusion Lipid-soluble Osmosis	substances enter and leave a cell by crossing substances can cross a cell surface membrane	its cell surface membrane
Describe how and Diffusion Lipid-soluble Osmosis	substances enter and leave a cell by crossing substances can cross a cell surface membrane	its cell surface membrane
Describe how and Diffusion Lipid-soluble Osmosis	substances enter and leave a cell by crossing substances can cross a cell surface membrane	its cell surface membrane

Oxygen and chloride ions can diffuse across cell-surface membranes. The diffusion of chloride ions involves a membrane protein. The diffusion of oxygen does not involve a

(b)

membrane protein.

A	Alveoli													
С	Capillaries												_	
D	Diffusion												_	
٧	entilation/												_	
													_	
_													_	
													_	
													_	
_													_	
_													_	
												(To	tal 10	m
a) D	Describe the one of the one of the out.	gross s	tructu	re of th	ne huma	an gas (exchan	ge systei	m and	d how	v we	breat	he in	
a) D a	Describe the good out.	gross s	tructu	re of th	ne huma	an gas (exchan	ge syste	m and	d how	we	breat	he in	
a) D	ind out.	gross s	tructu	re of th	ne huma	an gas (exchan	ge syste	m and	d how	v we	breat	he in	
a) D	Trachea	gross s	tructu	re of th	ne huma	an gas (exchan	ge syste	m and	d how	v we	breat	he in - -	
a) D	Trachea	gross s	tructu	re of th	ne huma	an gas (exchan	ge syster	m and	i how	v we	breat	he in - - -	
a) D	Trachea Bronchi Bronchioles	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	d how	v we	breat	he in	
a) D	Trachea Bronchi Bronchioles Alveoli	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	d how	v we	breat	he in	
a) D	Trachea Bronchi Bronchioles Alveoli Intercostal	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	d how	v we	breat	he in	
a) D	Trachea Bronchi Bronchioles Alveoli Intercostal muscles	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	d how	v we	breat	he in	
a) D	Trachea Bronchi Bronchioles Alveoli Intercostal muscles Diaphragm	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	d how	v we	breat	he in	
a) D	Trachea Bronchi Bronchioles Alveoli Intercostal muscles Diaphragm Volume /	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	d how	v we	breat	he in	
a) Da	Trachea Bronchi Bronchioles Alveoli Intercostal muscles Diaphragm Volume /	gross s	tructu	re of th	ne huma	an gas o	exchan	ge syster	m and	i how	v we	breat	he in	

Describe and explain how the lungs are adapted to allow rapid exchange of oxygen between air in the alveoli and blood in the capillaries around them.

Bond type?		
Saturation		
Solubility -		
Fatty acids		
Mucus also co	ontains glycoproteins. One of these glycoproteins is a polypeptide with , attached.	the
sugar, lactose	, attached.	the
sugar, lactose Describe how		the
sugar, lactose Describe how polypeptide to	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to Condensation	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to Condensation Bond type?	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to Condensation Bond type?	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to Condensation Bond type?	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to Condensation Bond type?	, attached. lactose is formed and where in the cell it would be attached to a	the
sugar, lactose Describe how polypeptide to Condensation Bond type?	, attached. lactose is formed and where in the cell it would be attached to a	the

Q18.

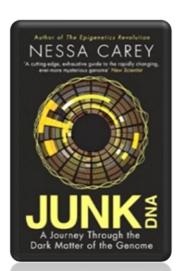
Lumen			<u>—</u>
Thickness			
Permeability			
Diffusion			<u> </u>
			_
Explain how tissue fluid is formed ar	d how it may be retu	urned to the circulatory sys	stem.
	•		
rterial end			
ressure			
Vater potential			
Osmosis			
vmph svstem			_
			_
			_
			_ _ _

Q19.(a) Explain two ways in which the structure of fish gills is adapted for efficient gas exchange:	
1	
2	
(b) Explain how the counter-current mechanism in fish gills ensures the maximum amount of oxygonsses into the blood flowing through the gills.	en
Lamellae	
Filaments	
Diffusion	
opposite	
Q20.	
The cardiac cycle is controlled by the sinoatrial node (SAN) and the atrioventricular node (AVN).
Describe how.	
SAN	
AVN	
Bundle of His	
Purkyne fibres	

(Total 5 marks)

Additional Reading/ Supporting Resources

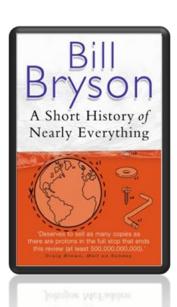


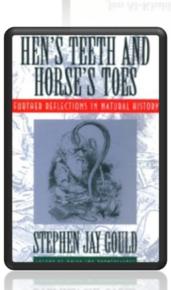


Junk DNA

Our DNA is so much more complex than you probably realize, this book will really deepen your understanding of all the work you will do on Genetics. Available at amazon.co.uk

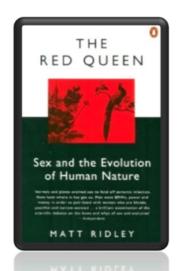
Studying Geography as well? Hen's teeth and horses toes Stephen Jay Gould is a great Evolution writer and this book discusses lots of fascinating stories about Geology and evolution. Available at amazon.co.uk





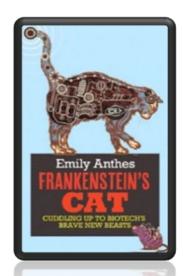
The Red Queen

Its all about sex. Or sexual selection at least. This book will really help your understanding of evolution and particularly the fascinating role of sex in evolution. Available at amazon.co.uk



A Short History of Nearly Everything

A whistle-stop tour through many aspects of history from the Big Bang to now. This is a really accessible read that will re-familiarise you with common concepts and introduce you to some of the more colourful characters from the history of science! Available at amazon.co.uk



An easy read.. Frankenstein's cat

Discover how glow in the dark fish are made and more great Biotechnology breakthroughs. Available at amazon.co.uk

Additional Reading Log:

Additional Reading Title	Dates?	Points of interest	How does it link to the course?