Candidate Name:

2023 End-of-Year Examination

Pre-University 3

H2 Biology

Paper 3 Long Structured and Free-response Questions

READ THESE INSTRUCTIONS FIRST

Do not open this booklet until you are told to do so.

Write your Admission number and name on all the work you hand in. Write in dark blue or black pen on both sides of the paper. You may use an HB pencil for any diagrams or graphs. Do not use staples, paper clips, highlighters, glue or correction fluid.

Section A

Answer **all** questions in the spaces provided on the Question Paper.

Section B

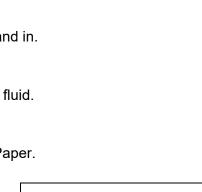
Answer any **one** question in the spaces provided on the Question Paper.

The use of an approved scientific calculator is expected, where appropriate.

You may lose marks if you do not show your working or if you do not use appropriate units.

The number of marks is given in brackets [] at the end of each question or part question.

For Examiner's Use		
Section	Question	Marks
	1	/30
Α	2	/10
	3	/10
В	4/5	/25
Total	/75	





Class

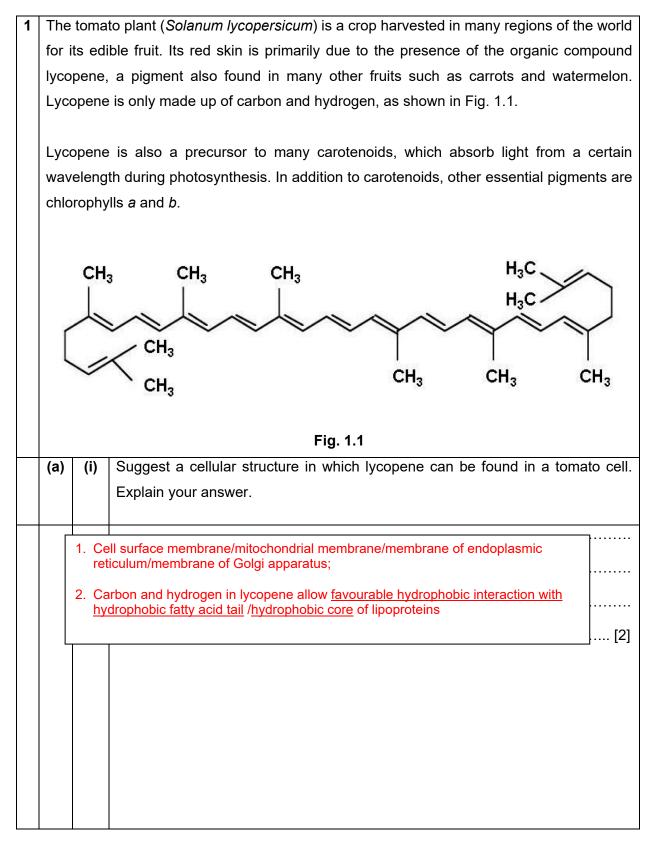
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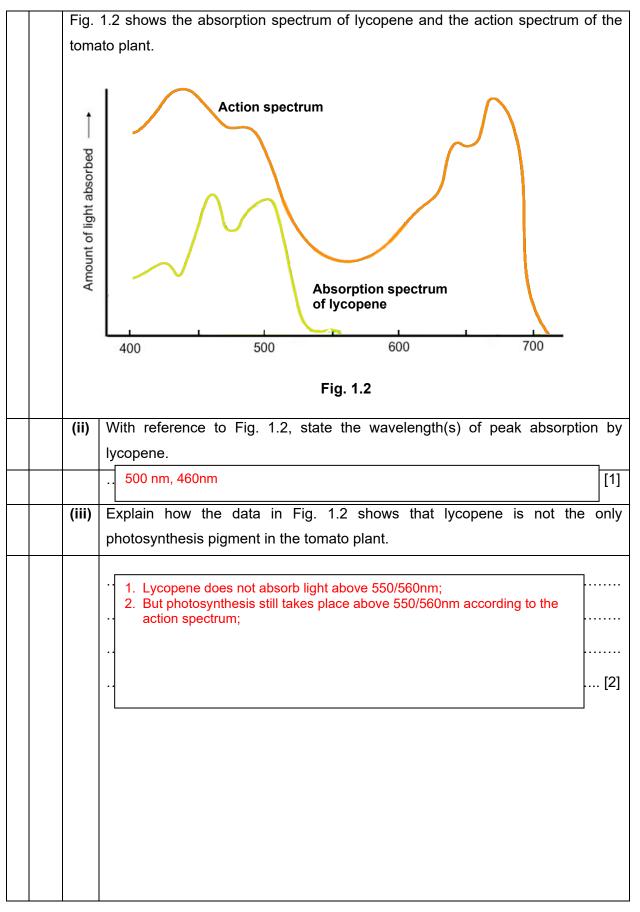
20 September 2023

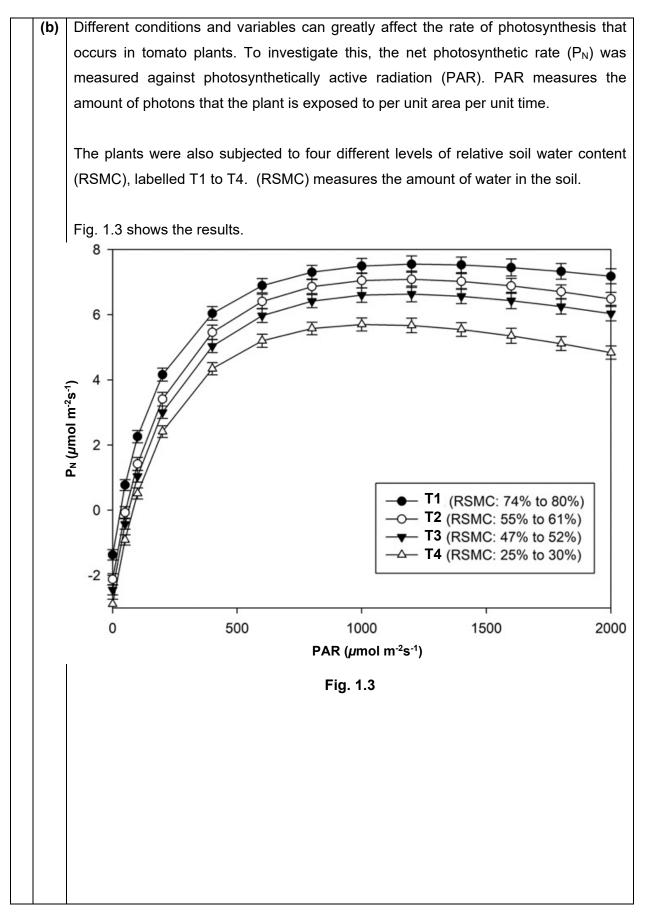
2 hours

Section A

Answer **all** questions in this section.







4

(i)	Explain the overall relationship between P_N and RSMC from 0 to 1000 μ mol μ	m⁻
	² S ⁻¹ .	_
	 As RSMC increases from T4, to T1, the highest P_N value increases from 5.8 <u>µmol m⁻²s⁻¹ to 7.8 µmol m⁻²s⁻¹;</u> Higher RSMC means there is a higher water content, which also means there is a higher rate of photolysis of water; Allows for more electrons to be passed down the electron transport chain during light dependent reaction, increasing production of <u>ATP and NADPH</u>; ATP and NADPH used for <u>Calvin Cycle to form G3P</u>, thus increasing the rate of photosynthesis; 	
(ii)	Suggest and explain the overall trend observed from 1500 to 2000 μ mol m- 2 s- $^-$	¹ .
2. <u>P</u>	he $\underline{P_N \text{ of all curves}}$ gradually decreases from 1500 to 2000 µmol m ⁻² s ⁻¹ ; AR/light intensity is no longer a limiting factor; ate of photosynthesis decrease due to heat stress;	
(iii)	Using Fig. 1.3, estimate the P_N when the PAR is 750 μ mol m ⁻² s ⁻¹ and the RSN is between 55% to 61%.	1C
	<u>6.5-6.8 μmol m⁻²s-1</u>	[1]

	(c)	The products of the light-dependent reaction are important for the form	ation of
		carbohydrates in plants.	
		Outline how carbohydrate can be continually formed using these products.	
			1
		1. <u>Carbon dioxide reacts with RuBP, cataysed by Rubisco to eventually form</u>	
		2. <u>ATP and NADPH reduce PGA to form G3P;</u>	
		3. For every <u>3 molecules of CO2</u> , there are <u>6 molecules of G3P</u> ;	
		4. <u>One molecule of G3P</u> exits the cycle to be <u>used as carbohydrate;</u> 5. the other 5 molecules form PuPP using ATP/PuPP is regenerated and can	
		5. the <u>other 5 molecules form RuBP using ATP/RuBP is regenerated</u> and can continue to <u>fix carbon dioxide;</u>	
		Any 4 points for 4 marks;	
			[4]
L	I		

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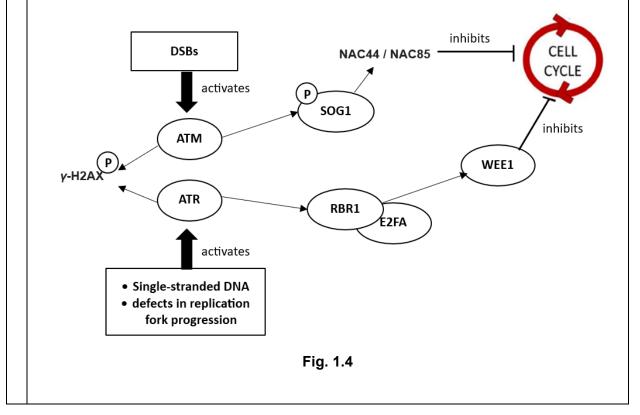
Stressful conditions can lead to the damaging of plant DNA. This leads to the activation of multiple cell signalling pathways to repair the DNA, which involves halting the cell cycle at its checkpoints.

Fig. 1.4 shows one such simplified pathway, known as the DNA damage response (DDR) signalling pathway. DDR activation depends on two protein kinases, ATM and ATR. ATM responds to double-stranded breaks (DSBs), where a double-stranded DNA breaks off at both strands, and ATR is activated by the presence of single-stranded DNA and any defects in replication fork progression. The result of the pathway is the inhibition of cell cycle progression.

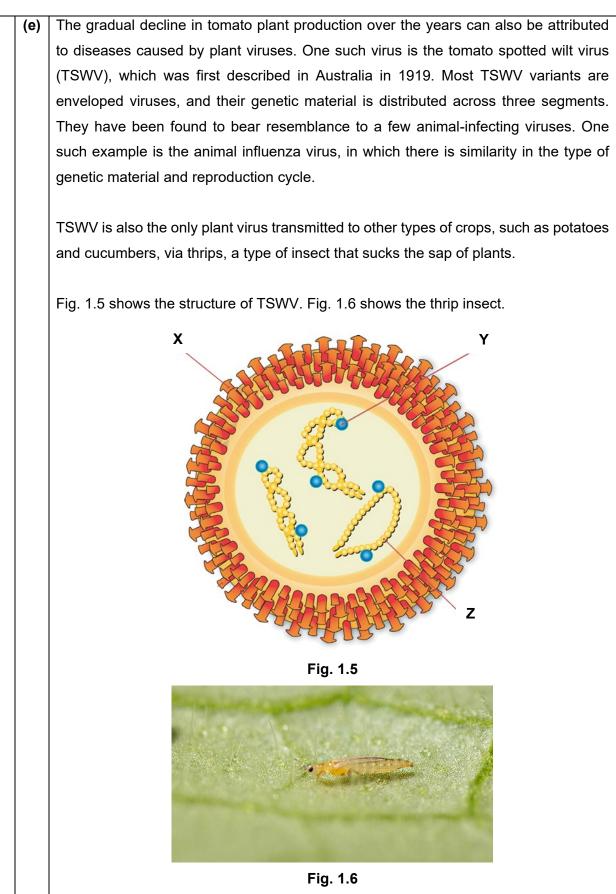
In response to the activation of ATM and/or ATR, the following pathways occur:

- Phosphorylation of the histone protein H2AX to form γ -H2AX (gamma-H2AX).
- The activation of SOG1, a transcription factor, activating proteins NAC44 and NAC85, which in turn will lead to the regulation of the M checkpoint.
- The expression of RBR1 and E2FA proteins, which are tumour suppressor proteins and WEE1, which is a G2 Checkpoint Kinase.

Through the activation of DDR, the DNA is subsequently repaired.



	Whi	ch protein k	inase ATM	Table 1.1 Protein(s)	Processes due to activation of
á			e activated?	involved	ATM / ATR
-	ATM a	nd ATR	H2AX		formation of heterochromatin
	ATR ATM		RBR1, E2FA, V SOG1, NAC44		cell size stops increasing
		for each row		, 14000	spindle fibres not attached to centromere
(d)	(i)	Complete	e Table 1.1 by fi	lling in the followi	ng:
		• If pro	otein kinase AT	M, ATR or both a	ctivated the pathway.
		• The	protein(s) that f	fit the description:	H2AX, SOG1, NAC44, NAC85, RBF
		E2F	A and WEE1. `	You are to fill in a	all the protein names in the table, a
		vou	only need to fill	l in each protein c	
		,	,	i în odon protoin c	106.
			,		
	(ii)	Explain th	-	of inhibiting cell	
		Explain the DSBs and	he importance d single-strande	of inhibiting cell ed DNA.	
		Explain the DSBs and 1.To preven of cancer;	he importance d single-strande it <u>accumulation c</u>	of inhibiting cell ed DNA. of DSBs and ssDNA	cycle progression in the presence which may lead to the <u>development</u> .
		Explain the DSBs and 1.To preven of cancer; 2. If this occonnected on the one of cancer on the one of the one	he importance d single-strande it <u>accumulation c</u> surs, this may lea <u>s</u> and <u>loss of fun</u>	of inhibiting cell ed DNA. <u>of DSBs and ssDNA</u> id to <u>gain of functio</u> ction mutation of tu	cycle progression in the presence which may lead to the <u>development</u> .
		Explain the DSBs and 1.To preven of cancer; 2. If this occord oncogene 3.To also prevention of the pre	he importance d single-strande at <u>accumulation o</u> surs, this may lea <u>s</u> and <u>loss of fun</u> event passing of	of inhibiting cell ed DNA. <u>If DSBs and ssDNA</u> id to <u>gain of functio</u> <u>ction mutation of tu</u> <u>mutations to daug</u>	cycle progression in the presence which may lead to the <u>development</u> .
		Explain the DSBs and 1.To preven of cancer; 2. If this occord oncogene 3.To also prevention of the pre	he importance d single-strande at <u>accumulation o</u> surs, this may lea <u>s</u> and <u>loss of fun</u> event passing of	of inhibiting cell ed DNA. <u>If DSBs and ssDNA</u> id to <u>gain of functio</u> <u>ction mutation of tu</u> <u>mutations to daug</u>	cycle progression in the presence which may lead to the <u>development</u> <u>n mutation of proto-oncogenes to</u> <u>mour suppressor genes;</u> <u>hter cells;</u>
		Explain the DSBs and 1.To preven of cancer; 2. If this occord oncogene 3.To also prevention of the pre	he importance d single-strande at <u>accumulation o</u> surs, this may lea <u>s</u> and <u>loss of fun</u> event passing of	of inhibiting cell ed DNA. <u>If DSBs and ssDNA</u> id to <u>gain of functio</u> <u>ction mutation of tu</u> <u>mutations to daug</u>	cycle progression in the presence which may lead to the <u>development</u> <u>n mutation of proto-oncogenes to</u> <u>mour suppressor genes;</u> <u>hter cells;</u>
		Explain the DSBs and 1.To preven of cancer; 2. If this occord oncogene 3.To also prevention of the pre	he importance d single-strande at <u>accumulation o</u> surs, this may lea <u>s</u> and <u>loss of fun</u> event passing of	of inhibiting cell ed DNA. <u>If DSBs and ssDNA</u> id to <u>gain of functio</u> <u>ction mutation of tu</u> <u>mutations to daug</u>	cycle progression in the presence which may lead to the <u>development</u> <u>n mutation of proto-oncogenes to</u> <u>mour suppressor genes;</u> <u>hter cells;</u> should there be any damage;
		Explain the DSBs and	he importance d single-strande t <u>accumulation o</u> trurs, this may lea <u>s</u> and <u>loss of fun</u> event passing of me for cell machi	of inhibiting cell ed DNA. of DSBs and ssDNA of to gain of function ction mutation of tu f mutations to daug inery to be repaired	cycle progression in the presence which may lead to the <u>development</u> <u>n mutation of proto-oncogenes to</u> <u>mour suppressor genes;</u> <u>hter cells;</u>



9744/03/PU3 EOY/2023

Using the information provided and your own knowledge of the influenza virus,		
(i)	Identify the structures X, Y and Z. X: Y: Z:	X: glycoproteins Y: RNA dependent <u>RNA polymerase</u> / <u>nucleoprotein</u> Z: negative sense single-stranded <u>RNA</u> [3]
(ii)	Describe how the genetic materia reproduction cycle.	I is replicated and used in the TSWV
	<u>capsid proteins</u> and <u>viral enzymes</u> <u>viral glycoproteins;</u>	mentary positive sense RNA; ate to synthesise negative sense A to be translated by host protein- bosomes in the cytosol to synthesise and bound ribosomes to synthesise
(iii)	- · ·	nit the virus from one plant to another.
	Acts as a vector/can carry the virus to undergo extrinsic incub another plant	ation period before transmitting to

[Total: 30 marks]

2	Dise	eases	can be characterised by defects that occur in many protein structures	s. For
	exa	mple,	there are several diseases related to defects in the haemoglobin structure,	which
	can	lead o	conditions such as anaemia.	
	(a)	State	e the effect on the haemoglobin's structure for the following defects and	how it
	(*)		ts the function of the haemoglobin.	
		(i)	No iron found in haem group	
				7
			fect: oxygen unable to <u>reversibly bind to haem group</u>	
			fected function: prevent uptake and release of oxygen/affect cooperativity binding	
			<u>oxygen</u>	
				[2]
	_	(ii)	Hydrophilic amino acid residues in the interior of structure.	_
			t: no deep hydrophobic cleft;	
		Aπec OR	ted function: no hydrophobic environment for binding of haem group	
			t: <u>hydrophobic amino acid residues</u> on the <u>exterior</u> of haemoglobin;	
		OR	ted function: haemoglobin is no <u>longer soluble in aqueous environment;</u>	
			t: loss of specific <u>3D conformation of haemoglobin;</u> ted function: haemoglobin no longer able to bind to oxygen;	
		Allee	ted function. Haemoglobin no longer able to bind to oxygen,	[2]
	Th€			such
	dise			prder
	whe	reвo	cells are unable to mature and the individual s immunity is compromised.	
	(b)	(i)	Describe three structural features of immunoglobin which allow it to perfo	orm its
			function.	
	1.		h immunoglobulin has two identical antigen binding sites that are	
	2.		<u>plementary to the epitope</u> determines <u>antigen binding specificity;</u> two identical antigen binding sites to bind to two antigens at the same time;	
	3.	<u>Con</u>	stant region determines its class/effector function/isotype of the antibody;	
	4.		<u>hinge region</u> which is made up of disulfide linkages provides the antibody <u>with</u> <u>ater flexibility</u> to better bind to the antigens/move independently;	
				[3]
				r _ 1

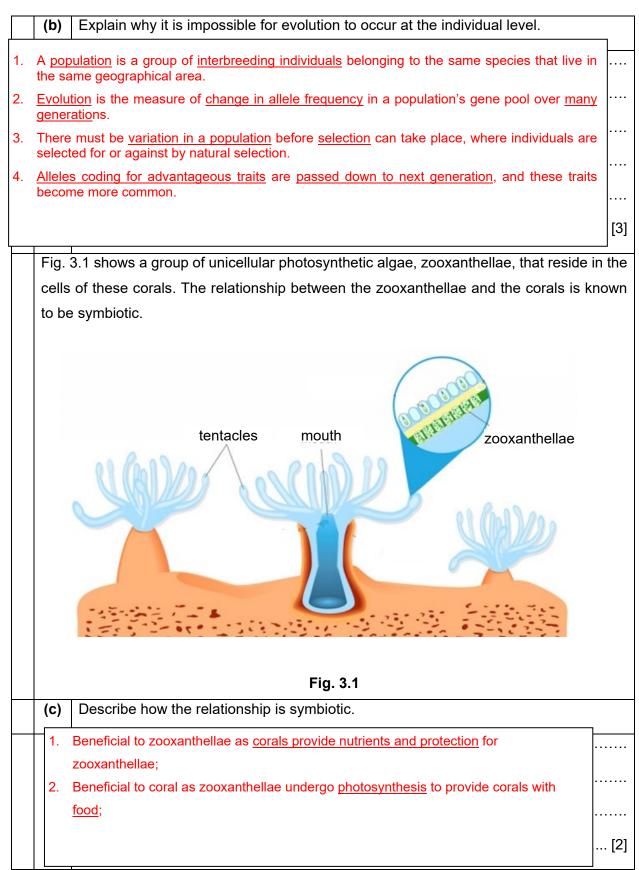
			(ii)	Explain why an individual's immunity is compromised when B cells are una mature.	able to
1. 2. 3.	<u>Tł</u> b c c	nere w a. <u>Ne</u> b. Bir c. Ac d. Co <u>tox</u>	<u>vill be restratis</u> nd to <u>fo</u> tivate pat <u>tum</u> <u>kicity;</u>	differentiate into a plasma cell and <u>secrete antibodies/ immunoglobulins;</u> no antibodies that can (any one): se foreign antigens and prevent them from entering host cell; preign antigens for opsonisation and allow phagocytosis; complement system for <u>enhanced phagocytosis/opsonisation;</u> nour cells for natural killer cells to recognise and allow <u>antibody-dependent cellular</u> emory B cells for stronger secondary immune response;	 [2]
			(iii)	Researchers have found a way to genetically engineer plants to pro- antibodies, known as plantibodies. Plantibodies can be extracted and in into humans to fight diseases. Suggest how using plantibodies could be advantageous over the m	jected
		1.	No nee	methods of producing antibodies in humans.	
		2.	No <u>late</u>	ency period when using plantibodies; al: 10	[1] marks]

3 *Acropora* is a genus of coral known for their size and hardness. They are also known to be the main contributors to the building of reefs. They contain numerous species and their phylogeny is well studied.

The number of single nucleotide polymorphisms (SNPs) shared between species can help to illustrate the relationship between these corals. SNPs are variations at a single position in a DNA sequence among individuals. A total of 1000 SNPs per species was investigated between three species, *A. hyacinthus, A. robusta* and *A. pichoni*. Table 3.1 shows the number of SNPs unique to each species when compared with one another.

	Tabl	е	3.	1
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		Comparison be	tween species	Number of unique SNPs	
		A. hyacinthus	A. robusta	44	
		A. robusta	A. pichoni	724	
		A. pichoni	A. hyacinthus	487	
(a)	Usir	ng the information pro	ovided,		
	(i)	calculate the number	er of shared SNPs betwee	en each species in the space	below.
	B	Between <i>A. hyacinthu</i>	s and <i>A. robusta</i> : 956		
	В	Between <i>A. robusta</i> ai	nd <i>A. pichoni</i> : 276		
	Between A. pichoni and A. hyacinthus: 513				
			Between A. hyacinth	us and A. robusta:	
			Between A. robu	sta and A. pichoni:	
			Between A. pichoni and	A. hyacinthus:	[3]
	(ii)	determine which tw	o species have the most	recent common ancestor. E	Explain
		your answer.			-
	1.	Between A. hyacinthus	and <i>A. robusta</i> : (ecf)		
	2.	-	shared SNPs indicates that	t there is <u>a high number of</u>	
		-		wing they are the most closely	
		related;		· · ·	
					[2]
	Щ				



[Total: 10 marks]

Section B

Answer **one** question in this section.

Write your answers on the lined paper provided at the end of this Question Paper.

Your answers should be illustrated by large, clearly labelled diagrams, where appropriate.

Your answers must be in continuous pose, where appropriate.

Your answers must be set out in parts (a) and (b), as indicated in the question.

4	(a)	Prokaryotic and eukaryotic genomes are organised differently. Explain how these
		differences affect the regulation of gene expression in prokaryotes and eukaryotes.
		[10]
	(b)	With reference to the life cycle of <i>Aedes aegypti</i> , describe the impact of temperature
		on the mosquito vector and dengue transmission. [15]
		[Total: 25]
5	(a)	The cell theory is characterised by concepts that define a cell. Explain how these concepts can be accounted for by both mitosis and the different types of stem cells.
		[10]
	(b)	With reference to examples, explain how phenotype is linked to genotype and the
		environment. [15]
		[Total: 25]

END OF PAPER

4(a) Prokaryotic and eukaryotic genomes are organised differently. Explain how these differences affect the regulation of gene expression in prokaryotes and eukaryotes. [10]

Note: question is about how the differences affect the regulation, and not what the differences are. It should also be clear which genome has which structure even if not explicitly stated per point.

Chromatin level

- 1. Prokaryotic genome is <u>not associated with histones</u>, and thus <u>histone modification is not</u> <u>possible</u> at the chromatin level;
- 2. Eukaryotic genome has <u>DNA wound around histone proteins</u>, which allows for <u>histone</u> <u>acetylation and methylation</u>;
- 3. This also controls access of transcription machinery to DNA;
- 4. <u>Prokaryotic genome is smaller than eukaryotic genome</u>, thus <u>more genes coding</u> for more proteins in eukaryotic genome;

Operon

- 5. Prokaryotic genome <u>organised into operons/has polycistronic mRNA</u> where <u>many</u> <u>structural genes are under the control of one promoter;</u>
- 6. Allows for <u>more efficient/coordinated control of gene expression for prokaryotes</u> since genes <u>responsible for same function are clustered and expressed together</u>;
- 7. Eukaryotic genome has <u>monocistronic mRNA</u> where <u>only one gene is under the control of</u> <u>one promoter;</u>
- 8. Allows for individual control in eukaryotes but not in prokaryotes;

Transcriptional/Translational level

- 9. Eukaryotic genome has <u>different combinations of enhancers and silencers that allow for</u> <u>combinatorial and coordinate control</u> of genes;
- 10. Prokaryotic genome is mainly regulated by operator which switches the operon on and off;
- 11. Prokaryotic <u>genome is found in nucleoid region</u> but eukaryotic genome is <u>enclosed in</u> <u>membraned bound nucleus</u>;
- 12. transcription and translation can <u>occur simultaneously in prokaryotes</u>, which is not possible in eukaryotes;

Presence of introns

- 13. Prokaryotic genome has <u>no introns which allows simultaneous transcription and</u> <u>translation</u> to occur;
- 14. Eukaryotic genome has <u>introns</u> and thus has to undergo <u>post transcriptional modification</u> <u>to remove introns</u> to form a continuous mRNA;
- 15. <u>Alternative splicing</u> is also possible in eukaryotes which allows <u>different proteins to be</u> <u>formed from the same mRNA</u>;

4(b) With reference to the life cycle of *Aedes aegypti*, describe the impact of temperature on the mosquito vector and dengue transmission. [15]

Relationship between Life Cycle of aedes and temperature

- 1. <u>Aedes aegypti</u> is the primary vector of <u>dengue virus;</u>
- 2. Mosquitoes are <u>cold blooded animals/ ectotherm</u> where their body temperature is affected by the <u>environment</u>;
- 3. The egg of the mosquito develops into larvae, then pupae, and finally adult mosquito;
- 4. <u>Embryonic development</u> is usually completed in 48 hours in a <u>warm</u> and humid environment, but may take weeks in cooler environment;
- 5. Increased temperature leads to shorter hatching time of egg into larvae;
- 6. <u>Larval development</u> occurs in four stages and is dependent on <u>temperature</u>, availability and density;

Effect of Temperature on mosquito vector

- Temperature greatly influences its <u>behaviour</u>, <u>development</u>, <u>reproduction</u> and <u>survival</u> (at least 2);
- 8. Increased temperature <u>increases the rate of enzymae catalysed reactions</u> which lead to <u>higher metabolic</u> rates;
- 9. As metabolic rates increase, the <u>development rates/life cycle of egg, larva and pupa</u> <u>shortens/hasten;</u>
- 10. At higher temperature, insects will mature, mate and <u>reproduce</u> in a shorter span of time than normal and consequently there is a greater capacity to <u>produce more offspring/eggs</u> during the transmission period;
- 11. At higher temperature, adult <u>female</u> mosquitoes <u>digest blood faster</u> and feed more frequently,
- 12. thus increasing transmission intensity as more humans are bitten;
- 13. Increased temperature leads to <u>increase water temperature</u>, <u>larvae</u> take shorter time to mature and consequently there is greater capacity to produce more offspring;
- 14. Shorten developmental time of mosquito <u>increases its survival rate</u> as <u>egg</u>, <u>larvae and</u> <u>pupae</u> are <u>less susceptible to predators</u>, <u>diseases and parasitism (any 1)</u>;
- 15. At temperature higher than 30°C, the survival rate of larvae and pupae drop/OWTTE;

Effect of Temperature on dengue transmission

- 16. Global warming leads to <u>temperatures in the sub-tropical regions becoming more optimal</u> <u>for both mosquitoes</u> and dengue virus;
- 17. Global warming can lead to <u>higher humidity</u> (in the sub-tropics), more stagnant water <u>breeding sites of water available for mosquitoes to lay eggs; OR</u>
- 18. Increase temperature may also lead to less stagnant water, reducing stagnant water <u>breeding sites</u> for adult mosquitoes to lay eggs;
- 19. Increased temperature leading to <u>increased precipitation/rainfall increases</u> the number and quality of <u>breeding sites for mosquitoes</u>;

- 20. Global warming will result in the spread of this disease beyond the tropics;
- 21. to higher altitudes and higher latitudes where it is cooler;
- 22. The <u>tropics</u> could become <u>too hot for the mosquito to survive</u>, as such, they might be a <u>decrease in incidence of dengue in the tropics</u>;
- 23. Increased temperature may lead to <u>warmer and shorter winters</u>, allowing <u>more mosquitoes</u> <u>to survive</u> during and through winter as they can be active for a longer period of time.
- 24. In warmer climate, dengue <u>virus</u> complete <u>extrinsic incubation</u> within the female adult mosquito <u>faster</u>, thereby increasing the proportion of infectious vector;

QWC 1 mark: Answer in structured and clear paragraph, with coherent link between temperature, life cycle of mosquito and dengue transmission;

Cell theory (3 marks)

- 1. States that the cell is the basic unit of structure and organisation in organisms;
- 2. All organisms are made up of one or more cells;
- 3. All cells come from pre-existing cells;

How mitosis supports cell theory (2 marks)

- 4. In humans, <u>genetically identical daughter nuclei</u> are formed from <u>parental nuclei via</u> <u>mitosis;</u>
- 5. Multicellular organisms are made up of more than one cell via mitosis;

How stem cells support cell theory (max 5 marks)

- 6. Stem cells are <u>unspecialised</u> and <u>undifferentiated</u> cells;
- 7. Stem cells can undergo <u>extensive proliferation through mitosis</u> to form a population;
- 8. And <u>differentiation to give rise to specialised</u> cell types;
- In multicellular organisms, there are different types of stem cells<u>: totipotent, pluripotent,</u> <u>multipotent and unipotent;</u>
- 10. Totipotent stem cells can differentiate into all types of cells;
- 11. Pluripotent stem cells can differentiate into cells from all three germ layers of embryo;
- 12. Multipotent stem cells can differentiate into cells within a closely related family;
- 13. Unipotent stem cells are differentiated cells with the ability to self-renew indefinitely;

How phenotype is linked to genotype (max 4 marks)

- 1. The genotype of an <u>organism is its genetic make-up</u>, which is the combination of alleles on the same gene on homologous chromosomes;
- 2. Phenotype is the <u>physical or physiological traits of an individual</u> which are <u>observable/measurable</u>;
- 3. As a result of gene expression and environmental factor;
- 4. Genes are transcribed into mRNA which is further translated into proteins;
- 5. Protein folds into specific 3D conformation, which determines the phenotypes;

Specific examples of genotype to phenotype (max 4 marks)

- 6. In a <u>heterozygote, allele that shows itself in phenotype is dominant allele</u>, and <u>recessive</u> <u>allele is not expressed</u>/dominant allele masks recessive allele;
- 7. In <u>incomplete dominance</u>, heterozygote exhibits phenotype that is an intermediate between the two homozygous forms;
- 8. In <u>codominance</u>, <u>both alleles have an equal effect on the phenotype</u> of the offspring/<u>equally expressed in phenotype</u> of heterozygote;
- 9. In <u>sex-linkage</u>, genes are found on the <u>sex chromosomes</u> and <u>males with only one X</u> <u>chromosome will express the phenotype</u> of the single allele present;
- 10. In <u>epistasis</u>, the <u>presence of one allele</u> of a particular gene <u>affects the expression of</u> <u>other genes</u> due to <u>gene interaction</u>;
- 11. In <u>continuous variation</u>, <u>polygenic inheritance</u> occurs where <u>multiple genes are involved</u> and there is an <u>additive effect of each gene</u>;

How phenotype is linked to environment

- 12. The phenotype of a characteristics is determined by the gene(s) controlling that particular characteristic but the <u>degree of expression of these genes</u> may be influenced by the <u>environment</u>;
- 13. Environmental effect exerts an <u>effect on phenotype determine by polygenes/presence of</u> <u>two or more genes;</u>

Temperature: Himalayan Rabbits

- 14. <u>Temperature</u> affects the coat colour of Himalayan rabbits;
- 15. Black fur is due to the pigment synthesised by an enzyme;
- 16. <u>At low temperature, the gene coding for the enzyme that produces the pigment is</u> <u>expressed, giving rise to black fur;</u>
- 17. <u>At high temperature, the gene coding for the enzyme that produces the pigment is not</u> <u>expressed, giving rise to white fur;</u>

Diet: Queen and Worker bees

- 18. <u>Queen and workers have the same amount of genetic material but they are phenotypically different;</u>
- 19. These phenotypic differences are due to the diet of the larvae;

- 20. After hatching, all the larvae are fed with royal jelly;
- 21. Larvae destined to be workers are switched to a diet consisting of honey and pollen, whereas those destined to be queen continue with royal jelly;
- 22. The <u>high protein content of royal jelly stimulates the formation and maturation of the</u> <u>female reproductive system</u> in the <u>queen bees;</u>
- 23. <u>Mutagens in the environment</u> result in <u>gene mutations</u> resulting in changes in phenotype;

QWC Answer in structured and clear paragraph, with at least 2 points from all three paragraphs.