

AS Level Biology A

H020/02 Depth in biology

Tuesday 7 June 2016 – Afternoon

Time allowed: 1 hour 30 minutes

You must have:

• the Insert (inserted)

You may use:

- · a scientific calculator
- a ruler (cm/mm)



First name	
Last name	
Centre number	Candidate number

INSTRUCTIONS

- Use black ink. You may use an HB pencil for graphs and diagrams.
- Complete the boxes above with your name, centre number and candidate number.
- · Answer all the questions.
- Write your answer to each question in the space provided. If additional space is required, you should use the lined page(s) at the end of this booklet. The question number(s) must be clearly shown.
- · Do not write in the barcodes.

INFORMATION

- The total mark for this paper is 70.
- The marks for each question are shown in brackets [].
- Quality of extended responses will be assessed in questions marked with an asterisk (*).
- · This document consists of 24 pages.

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Turn over

Answer **all** the questions.

1 (a) Fig. 1.1 is a diagram that represents inspiration and expiration in a human.

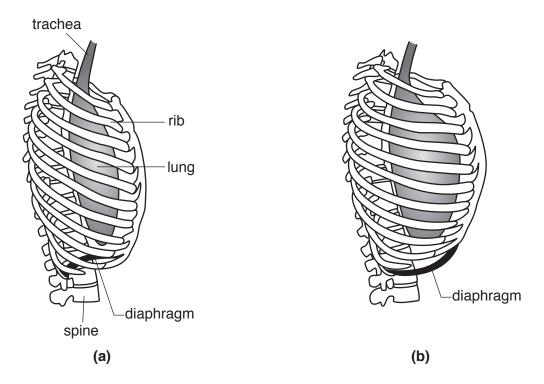


Fig. 1.1

(i)	Which of the two diagrams, (a) or (b), represents the body immediately after expiration	n?
	Describe how this diagram justifies your choice.	
		[2]
(ii)	Why can expiration be a passive process?	
		[1]

(iii)	Some chemicals can act as allergens. If these allergens are inhaled, they can cause breathing problems. Allergens cause the smooth muscle in the walls of the airways to contract.
	Suggest the effects that this muscle contraction has on ventilation.
	[6]

(b) Fig. 1.2 represents the volume changes in the lung of a human.

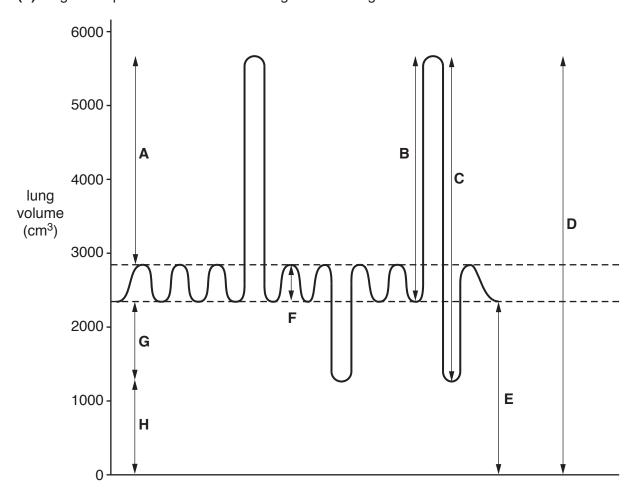


Fig. 1.2

(i) Select the letter, A to H, that corresponds to each of the following lung volumes.

The first one has been done for you.

Lung volume	Letter
Inspiratory reserve volume	Α
Residual volume	
Total lung capacity	
Tidal volume	
Vital capacity	

[4]

(ii)	Volume C can be measured using an instrument such as a spirometer.
	What breathing instructions would be given to a person whose volume C was being measured?
	[2]

2 (a) Mitosis and meiosis play an important role in the life cycles of organisms.

Fig. 2.1 and Fig. 2.2 represent an outline of the life cycles of two different organisms.

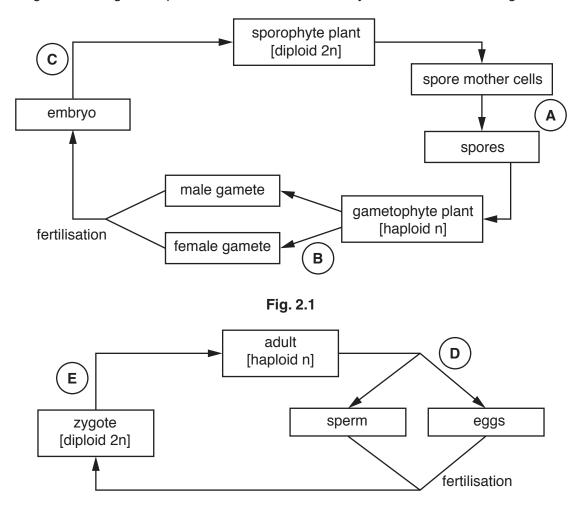


Fig. 2.2

Place a tick (\checkmark) in each row of the table to indicate the type of nuclear division that occurs at each of the letters **A** to **E**.

	Mitosis	Meiosis
Α		
В		
С		
	Mitosis	Meiosis
D	Mitosis	Meiosis

(b) Fig. 2.3 is a diagram that represents the different phases of the cell cycle.

X, Y and Z represent checkpoints in the control of the cell cycle.

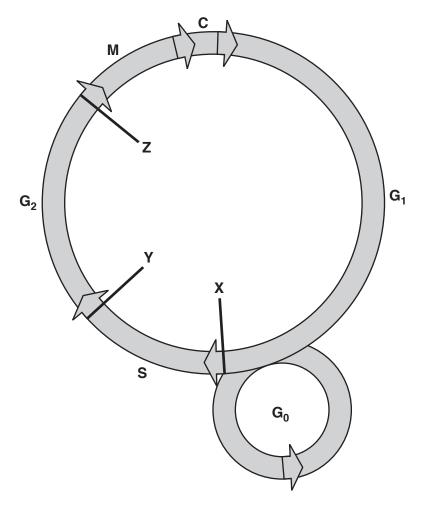


Fig. 2.3

(i)	State all the letters in Fig. 2.3 that represent the phases of interphase.
	[1]
(ii)	Suggest what is being checked at checkpoint Y on Fig. 2.3.
	[1]

(c) Table 2.1 indicates the relative time spent in different phases of the cell cycle for three different types of cell, P, Q and R.

Call type	Relative time spent in a phase				
Cell type	G ₁ /G ₀	S	G ₂	M/C	
Р	18	50	13	19	
Q	18	25	11	16	
R	100	0	0	0	

Table 2.1

(1)	which of the cells P , Q of R takes the shortest time to divide?
(ii)	Suggest why cell P spends twice as much time in phase S than cell Q .
	[1]
(iii)	What can be deduced about the behaviour of cell R ? Give reasons for your answer.
	[0]

- (d) An experiment was carried out where a student observed cells in different tissues under the microscope.
 - The cells were undergoing mitosis.
 - 200 cells were observed for each tissue.
 - The number of cells in each stage of mitosis was recorded.

The results are shown in Table 2.2.

Tissue	Nun	Total			
type	Prophase	Metaphase	Anaphase	Telophase	Total
V	65	55	7	73	200
W	85	59	6	50	200

Table 2.2

The student had expected that the results observed for tissue type ${\bf W}$ would not be significantly different from those for tissue type ${\bf V}$.

(i)	Identify the pieces of evidence in Table 2.2 that caused the student to suspect that the results for tissue type W might be significantly different from those for tissue type V .
	[1]
(ii)	The student decided to analyse the data using a statistical test.
	A friend suggested using Student's <i>t</i> -test.
	Why is Student's <i>t</i> -test not suitable for dealing with this data?
	[1]

- (e) The chi-squared (χ^2) test can be used to analyse the data.
 - (i) Complete the rows for metaphase and telophase in the table below and calculate the χ^2 value for the data.

The χ^2 value is calculated using the following formula:

$$\chi^2 = \sum \frac{(O - E)^2}{E}$$

Cells	Observed (O)	Expected (E)	(O-E)	(O-E) ²	(O – E) ² E
In prophase	85	65	20	400	6.154
In metaphase					
In anaphase	6	7	-1	1	0.143
In telophase					
Total	200	200			

(ii) The value of chi-squared (χ^2) can be used to conclude whether the results for cells in tissue type **W** differ significantly from those for tissue type **V**.

The number of **degrees of freedom** determines which row of the χ^2 probability table is used.

The number of degrees of freedom is defined as:

the number of categories - 1

What will be the number of degrees of freedom used in this analysis?

[1]

(iii) The student had expected that the results observed for tissue type **W** would not be significantly different from those for tissue type **V**.

Use your calculated value for χ^2 and the information from the χ^2 probability table below to conclude whether or not the results observed for tissue type ${\bf W}$ are significantly different from those for tissue type ${\bf V}$.

Degrees of		Probability (p)							
freedom	0.99	0.95	0.05	0.01	0.001				
1	0.00	0.00	3.84	6.64	10.83				
2	0.02	0.10	5.99	9.21	13.82				
3	0.11	0.35	7.82	11.35 13.28	16.27 18.47				
4	0.30	0.71	9.49						
5	0.55	1.15	11.07	15.09	20.52				
6	0.84	1.64	12.59	16.81	22.46				
7	1.24	2.17	14.07	18.48	24.32				

 	 	 	[2]

3 (a) Polymers are important molecules that have structural and functional roles in organisms.

Chitin is a polymer that is found in insects, where it forms a major part of the structure of the exoskeleton.

- Chitin is a macromolecule that is similar to a polysaccharide.
- Chitin is composed of molecules of N-acetylglucosamine, the structure of which is shown in Fig. 3.1 below.
- The monomers of N-acetylglucosamine join by 1–4 glycosidic bonds to form the chitin molecule.

Fig. 3.1

(i)	How does the composition of N-acetylglucosamine differ from the composition monosaccharide sugar?	
(ii)		
		[2]

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(iii)	Using your knowledge of the formation of structural polysaccharides, describ formation of the chitin molecule from its monomer and predict its structure.	e the
		[4]

(b) Fig. 3.2 is a photomicrograph of the trachea of a honeybee, *Apis mellifera*.

The trachea of this honeybee is infected with honeybee tracheal mites, *Acarapis woodi*. Some of these mites are labelled **M** on Fig. 3.2.

The trachea and tracheoles of insects have circular bands of chitin. One of these bands is labelled ${\bf C}$ on Fig. 3.2.

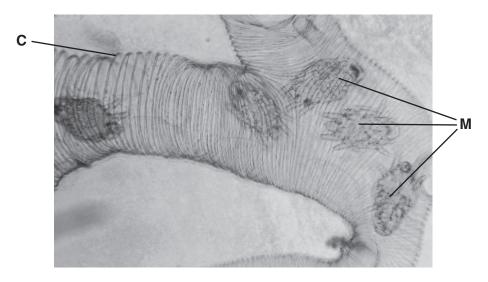


Fig. 3.2

(i)	What is the function of the circular bands of chitin labelled C?
	[1]
(ii)	The mites use their mouthparts to bite through the walls of the trachea. They then feed off the haemolymph, the blood-like liquid that bathes the cells and organs of the honeybee.
	Suggest one other way in which the presence of the mites might affect the honeybee.
	[1]

- **4 (a)** Fungi produce enzymes to digest complex food substances. Amylase is an enzyme that catalyses the conversion of starch to maltose.
 - A sample of the fungus Amanita citrina was placed on agar in a petri dish.
 - The agar contained starch.
 - The dish was incubated until the thread-like hyphae had grown a few centimetres.
 - lodine solution was then poured onto the surface of the agar.

A diagram representing the results is shown in Fig. 4.

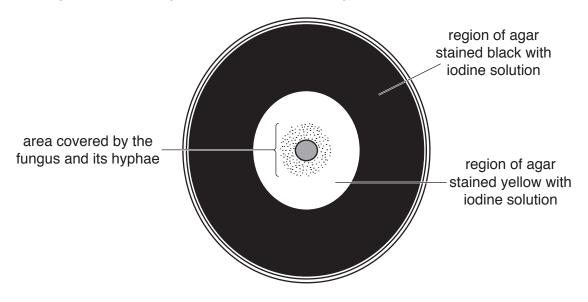


Fig. 4

(i)	To which genus does this fungus belong?
	[1]
(ii)	The region of yellow staining shown in Fig. 4 includes part of the agar where the fungus had not yet grown.
	What does this pattern indicate about the action of the fungal enzymes?
	[1]

(b) Lipase is an enzyme that catalyses the breakdown of lipids.

An investigation was carried out to see the effect of temperature on the activity of a lipase.

- 5 cm³ of an alkaline solution of lipid was poured into a test tube.
- The test tube was placed into a water bath maintained at 20 °C and left to equilibrate.
- A few drops of an indicator were added to the wells of a white spotting tile.
 The indicator is pink above pH values of 8.3 and turns colourless at pH values below 8.3.
- Once the lipid solution had equilibrated, 1 cm³ of 0.5% lipase solution at the same temperature was then added to the test tube.
- For five minutes, at 30 second intervals, the solution was stirred and a few drops were removed from the test tube and placed in a well on the white spotting tile.
- The time was recorded when the solution and indicator did not remain pink.
- The procedure was repeated four more times at 20 °C and then again at a further six temperatures.

The results are shown in Table 4.1 below.

Temperature	Time when solution did not remain pink								
(°C)	Replicate 1	Replicate 2	Replicate 3	Replicate 4	Replicate 5				
20	210 270		240	300	270				
25	25 90 120 30 60 60		210	180	120				
30			90	90	60				
35	60	60	60	90	60				
40	40 210 12 45 240 30		210	180	210				
45			300	_	270				
50	_	_	_	_	_				

Table 4.1

(i)	Why is pH not a controlled variable in this investigation?	
		[1]

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) I	dentify one variable that has been controlled in this procedure.
ı	dentify one variable, other than pH, that has not been controlled in this procedure.
	[1]
	The procedure required the solution to be stirred and then drops of solution to be placed on a white spotting tile.
	Suggest why this procedure was followed rather than simply adding indicator to the test tube, stirring the solution and looking for the colour change in the test tube.
	What can be concluded from the results in Table 4.1 about the optimum temperature for ipase activity?
	[1]
	Describe two different ways in which the procedure could be modified to obtain a more accurate value for the optimum temperature for lipase activity.
•	1
2	2
	[4]

t	There are two mo emperature can aff	ect these	mecha	inisms o	f lipase	action.	donom	- Cu		onang
									•••••	

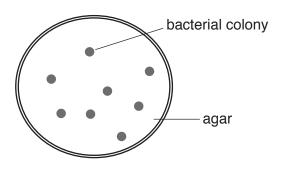
5

(a)	Fig.	5.1, on the insert , shows the circulatory systems of three groups of animals.
	(i)	What type of circulatory system is shown in all these animals?
		[1]
	(ii)	How does the circulatory system of a fish compare to that of a mammal?
		[1]
(b) ³	Fig.	5.2, on the insert , shows the flow of blood through the heart of an amphibian such as a l.
		e the information in Fig. 5.1 and Fig. 5.2 to compare the circulations of a frog and a mmal and the relative effectiveness of each type of circulation.
		[6]

6 (a) An experiment was carried out to investigate the resistance of a species of bacterium to the antibiotic penicillin.

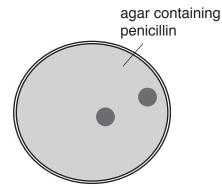
Bacteria were allowed to grow into colonies on an agar plate.

A cloth was placed onto the bacteria and then the pattern of bacterial colonies was transferred to an agar plate that contained penicillin.

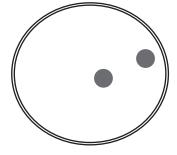


Only two colonies survived and continued to grow on the agar that contained penicillin.

The bacteria in these colonies possessed a mutation that gave them resistance to the penicillin.



The original plate was flooded with a solution containing penicillin and the same two colonies continued to grow.



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(i) A student made the following suggestion:

I think that the colonies on the agar containing penicillin that survived and grew did so because those bacteria evolved resistance.

They evolved resistance as a result of being exposed to the penicillin.

Another student commented:

But some of the bacteria in the population were already resistant, so they can't have evolved resistance because they were exposed to the penicillin.

	What evidence population?	indicates	that th	е р	enicillin-resistant		already			
										[1]
(ii)	Name the proce population.	ess that ir	ocrease	s the	e proportion of p	enicillin-r	esistant	bacteria	in	the
										[1]

(b) One role of the Office for National Statistics (ONS) is to collate data about the causes of death in England and Wales. Deaths involving *Staphylococcus aureus* and MRSA statistics have been produced by the ONS for each year since 1993.

S. aureus can be mentioned on a death certificate and *S. aureus* may also be specified as being methicillin resistant (MRSA).

Table 6 shows the data for the years 1993 to 2012.

	Number of death certificates mentioning S. aureus			
Year	S. aureus not specified as resistant	S. aureus specified as MRSA	Total	
1993	379	51	430	
1994	358	90	448	
1995	409	198	607	
1996	445	298	743	
1997	395	386	781	
1998	451	409	860	
1999	484	480	964	
2000	476	666	1036	
2001	473	731	1204	
2002	421	794	1215	
2003	448	968	1516	
2004	461	1138	1599	
2005	450	1649	2099	
2006	498	1652	2150	
2007	459	1593	2052	
2008	270	1230	1500	
2009	472	781	1253	
2010	475	485	960	
2011	274	364	638	
2012	265	292	557	

Table 6

MRSA from 1993 to the year when the numbers reach a peak.

(i) Calculate the percentage increase in the number of death certificates that mention

	Show your working and give your answer to three significant figures.
	Answer = % [2]
/ii\	
(ii)	The proportion of death certificates that mention MRSA in 1993 is 12%.
	Compare this figure with the proportion of death certificates that mention MRSA in 2012.
	[2]
(iii)	What can you conclude from these data about the deaths involving <i>S. aureus</i> and MRSA since 2007?
	[2]

END OF QUESTION PAPER

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ADDITIONAL ANSWER SPACE

	space is required, you should use the following lined page(s). The question number(sarly shown in the margin.
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