Centre Number	Candidate Number	Name

## UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Subsidiary Level and Advanced Level

**BIOLOGY** 9700/31

Paper 31 Advanced Practical Skills

For Examination from 2007

Specimen Paper

2 hours

Candidates answer on the Question Paper. Additional Materials: As listed in Instructions to Supervisors.

#### **READ THESE INSTRUCTIONS FIRST**

Write your Centre number, candidate number and name on all the work you hand in. Write in dark blue or black pen.

Do not use staples, paper clips, highlighters, glue or correction fluid.

You may use a pencil for any diagrams, graphs or rough working.

Answer all questions.

At the end of the examination, fasten all your work securely together.

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

1 You are reminded that you have only one hour for this part of the practical examination. You should read carefully through the whole of this question and then plan your use of the time to make sure that you finish all the work that you would like to do.

Respiration is a process which uses enzymes to release energy from biological molecules.

(a) You should spend no longer than five minutes on question 1 (a).

You are provided with a solution of a biological molecule, **\$1**. You are provided with the following materials that can be used to identify the biological molecule in solution **\$1**.

- Ethanol
- Benedict's solution
- Distilled water

Use the materials provided to identify the biological molecule in solution **S1**.

Describe each test that you performed and explain the meaning of the results that yo obtained.	u
	••
	••
[2	2]

(b) You should spend no more than 25 minutes on question 1 (b)

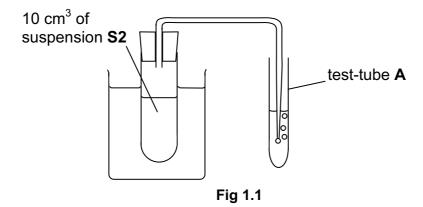
You are provided with a suspension of yeast that has been placed in solution S1, labelled **S2**.

Carefully follow the instructions below to use **S2** to investigate the quantitative effect of temperature on the enzymes in the yeast. You should present and record your observations and data in the space provided.

You will need to:

- read through the instructions carefully,
- make some decisions,
- prepare the space on the next page so that it is ready for you to record the readings.

- Place 10 cm<sup>3</sup> of suspension S2 into the large test-tube.
- Securely fit the bung with the delivery tube into the top of the large test-tube
- Place 5 cm<sup>3</sup> of distilled water in an empty test-tube, A.
- Place 150 cm<sup>3</sup> of water in the beaker.
- Measure the temperature of the water.
- Use the beaker as a water bath for the large test-tube, so that the delivery tube is outside the beaker
- Place test-tube **A** so that the end of the delivery tube is near the bottom of the water in test-tube **A**, as shown in Fig 1.1.



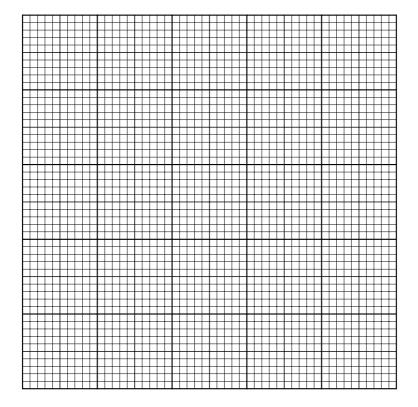
Bubbles of gas should come from the end of the delivery tube.

- (i) Decide how many readings to take, and for how long to take each reading. Each reading should be made and recorded in the space you have prepared below. [1]
- (ii) You can use the Bunsen burner to warm up the water in the water bath. Decide how many different temperatures you will use, and what would be appropriate temperatures to use. Repeat the readings taken in (ii) at each of your chosen temperatures.

[6]

(c)	(i)	pH has a big e was controlled			ne reactions.	Suggest how effectively pH
						[1]
	(ii)	State <b>two</b> sign	ificant source	es of error in t	his experimen	t, other than control of pH.
(d)		student's inves				[1] produced in five minutes was
				Table 1.1		
	tem	perature/°C	bubbles of gas produced ir minutes		ed in 5	mean number of bubbles of gas/
			first run	second run	run	bubbles min
	28		10	8	11	1.9
	38		19	21	17	3.8
	48		24	21	28	
	58		11	6	10	1.8
_	(i)	The first time to in five minutes repeated it.	that the studes was 3 cm <sup>3</sup>	ent tried this a	at 38 °C, the r ny the student	number of bubbles produced discarded the reading and
						[1]
	<b>(::</b> \	Ol-t- T-b				
	(11)	bubbles produ	•	•	nissing value i	for the mean number of gas
		Show your wor	rking in the s	pace below		

(iii) Plot a graph to show the effect of temperature on the mean number of gas bubbles produced, using the data in Table 1.1.



ı	101
	[၁]

(e)	Briefly outline the main features of the relationship between temperature and volume of gas produced.	mean
		[1]

	as the temperature increases, the rate of production of gas will increase	
	Draw an appropriate conclusion to the student's experiment, including	
	<ul> <li>whether the experimental data supports the student's hypothesis,</li> <li>a revised or new prediction.</li> </ul>	
		······
		[2]
(g)	Suggest how the experiment in question 1 (a) could be improved.	
		[2]

2	You	sho	reminded that you have only one hour for this part of the practical examination. buld read carefully through the whole of this question and then plan your use of the make sure that you finish all the work that you would like to do.
	(a)	(i)	Draw a low power plan diagram of the specimen on slide <b>\$4</b>
			roj.
		(::\	[2]
		(11)	Use a ruler to measure the actual size of the specimen on slide S4 and the size of your drawing across the same point. Draw a line on your drawing to show the size that you have measured. Calculate the magnification of your drawing.
			Show your working
			magnification[2]
		(iii)	Estimate the uncertainty in the measurement of the actual size of the specimen on
		(,	slide <b>S4</b> .
			A chual cima ne a a coma d —
			Actual size measured =
			Uncertainty =[1]

(iv) Suggest how a systematic error in measuring the size could occur.
7.41
[1]
(b) Starch is stored as granules in some of the cells in the specimen on slide <b>S4</b> . Starch is stained purple during preparation of slide <b>S4</b> .

In the space below, show your observations of enough of these food storage cells to give a representative sample of the range of their structure.

[6]

(c)	(i)	Prepare the space below so that it is suitable for you to compare, using a hand lens and microscope, specimen <b>S3</b> and the specimen on slide <b>S4</b> .
		[1]
	(ii)	Compare specimen <b>S3</b> and the specimen on slide <b>S4</b> , recording your observations in the space that you prepared in question <b>2 (c) (i)</b> . [3]
	(iii)	Both specimens are involved in transport of materials. State one key point of your observations that relates to this function.
		[1]

(d) The photomicrographs, Fig. 2.1 and Fig. 2.2 are taken from a different part of an unfamiliar plant. Fig. 2.1 is a transverse section across the structure, and Fig. 2.2 is a longitudinal section along it.

Mature xylem vessels are large tubes with thick cell walls and no cytoplasm within them. As they mature the cells die, the end walls of the cells break down and they become a continuous tube. They are found within vascular bundles that run along roots, stems and inside the veins of leaves. They do not have companion cells. The cell walls of the xylem vessels or the cells next to them may have rings or spirals of thickening, and may have pits, which are holes through the cell walls connecting cells with the cell next to them.

Use clear labels and label lines to show the xylem in each of the photomicrographs using the information provided. Explain the reasons for your choice in the space provided.

	A TOTAL
Reasons for choice	

Fig. 2.1

محال الله
THE RESERVE OF THE PARTY OF THE

Fig. 2.2

[2]

© UCLES 2006 9700/31/SP07

Reasons for choice

#### **BLANK PAGE**

Copyright Acknowledgements:

 $\label{eq:Question 2} \textbf{Question 2} \qquad \qquad \textbf{Fig. 2.2} \ @ \ \textbf{http://www.lima.ohio-state.edu/academics/biology/images/cornstemls.jpg}$ 

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Centre Number	Candidate Number	Name

# UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Subsidiary Level and Advanced Level

BIOLOGY 9700/31

Paper 31 Advanced Practical Skills

For Examination from 2007

SPECIMEN INSTRUCTIONS

Great care should be taken that any information given does not reach the candidates either directly or indirectly.

This document consists of 4 printed pages.

#### Instructions for preparing apparatus

These instructions give details of the apparatus and materials required by each candidate for this paper. Sufficient information is given to permit the Centre to set up and test the apparatus and materials so that the candidates can be fairly assessed.

If a candidate breaks any of the apparatus, or loses any of the material supplied, the matter should be rectified.

Candidates must be provided with a microscope with:

- Low-power objective lens, e.g. X10 (equal to 16 mm or <sup>2</sup>/<sub>3</sub>")
- High-power objective lens, e.g. X40 (equal to 4 mm or ½")
- Eyepiece graticule fitted within the eyepiece and visible in focus at the same time as the specimen.

Each candidate should have sole, uninterrupted, use of the microscope for at least 55 minutes.

Supervisors are advised to remind all candidates that **all** substances in the examination should be treated with caution. Pipette fillers and safety goggles should be used when necessary.

In accordance with the COSHH (Control of Substances Hazardous to Health) Regulation, operative in the UK, a hazard appraisal of the examination has been carried out.

The following codes are used where relevant.

**C** = corrosive substance

H = harmful or irritating substanceF = highly flammable substance

**O** = oxidising substance

T = toxic substance

Centres are reminded that they are **not** permitted to open any question paper envelopes before the examination. Centres are also referred to the Handbook for Centres 2007, and in particular Section 3.1.2 (c), Security of Question Papers and Examination Materials, as well as 3.3.11.1, Practical Examinations in Science Subjects.

#### **Instructions to Supervisors**

Each candidate must be provided with the following apparatus and materials.

#### To be supplied by the Centre

#### **Question 1**

Each candidate will require:

- (i) 20 cm<sup>3</sup> of solution S1, labelled **S1**. This should be a 0.5 mol dm<sup>-3</sup> glucose solution. It could be made by dissolving 9 g of glucose in 80 cm<sup>3</sup> of water and making up to 100 cm<sup>3</sup>.
- (ii) A small volume (e.g. 10 cm<sup>3</sup>) of Benedict's solution in a suitable dispensing bottle, labelled **Benedict's solution**.
- (iii) A small volume (e.g. 10 cm<sup>3</sup>) of ethanol or industrial methylated spirit ('meths') in a suitable dispensing bottle, labelled **ethanol**.
- (iv) At least 20 cm<sup>3</sup> of distilled water in a small dispensing bottle or a container with a pipette, labelled **distilled water**.
- (v) Test-tube rack containing two empty, unlabelled test-tubes, a large test-tube and a test-tube labelled **A**.
- (vi) Water-bath to perform Benedict's test, consisting of a Bunsen burner, tripod, gauze and beaker.
- (vii) A bung and glass or plastic delivery tube, as shown in Fig. 1.1, to fit the large test-tube in (v).

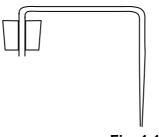


Fig. 1.1

- (viii) A beaker large enough to contain 150 cm<sup>3</sup> of water and not overflow when the large test-tube is dipped in the water. This could be the same beaker as is used for the waterbath in (vi).
- (ix) Access to a tap dispensing water at, or below, room temperature.
- (x) 20 cm<sup>3</sup> of yeast suspension S2, labelled **S2**. This should be made using a 1.0 mol dm<sup>-3</sup> glucose solution. It could be made by dissolving 18 g of glucose in 80 cm<sup>3</sup> of water. This can be made up several days before the examination. About half-an-hour before the examination, 1 g of dried yeast powder or granules should be thoroughly stirred into the glucose solution, and this should

- be made up to 100 cm<sup>3</sup> before the yeast starts to ferment and produce foam, which will make measurement of the volume difficult.
- (xi) Thermometer capable of measuring at least from 0 to 60 °C (e.g. a standard -10 to 110 °C laboratory thermometer would be suitable)
- (xii) A Bunsen burner. This could be the same burner as is used to heat the waterbath in (vi).

#### **Question 2**

Each candidate will require:

- (i) Specimen **S3**, a slice from a carrot, between 1 cm and 6 cm in diameter, and between 0.5 and 1 cm thick. The central stele should be clearly visible in the carrot used.
- (ii) Slide **S4**, a transverse section of *Ranunculus* root, suitably stained to show the central stele and starch grains in the parenchyma cells. (A suitable slide may be purchased from CIE, through the publications catalogue.)
- (iii) A microscope with:
  - Low-power objective lens, e.g. x10 (equal to 16 mm or <sup>2</sup>/<sub>3</sub>")
  - High-power objective lens, e.g. x40 (equal to 4 mm or  $\frac{1}{6}$  ")
  - Eyepiece graticule fitted within the eyepiece and visible in focus at the same time as the specimen.
- (iv) Hand lens (e.g. x10)

### **UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS**

Specimen for 2007

## **GCE A/AS LEVEL**

# MARK SCHEME

**MAXIMUM MARK: 40** 

SYLLABUS/COMPONENT: 9700/31

BIOLOGY ADVANCED PRACTICAL SKILLS



Page 2	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

Skill	Total marks	Breakdown of mark expect	ations	Question 1	Question 2
Manipulation, measurement	16 marks	Successful collection of data and observations	8 marks	2	6
and observation		Decisions relating to measurements or observations	8 marks	4	4
Presentation of data and observations	12 marks	Recording data and observations	4 marks	2	2
observations		Display of calculation and reasoning	2 marks	1	1
		Data layout	6 marks	4	2
Analysis, conclusions and	12 marks	Interpretation of data or observations and identifying sources of error	6 marks	2	4
evaluation		Drawing conclusions	3 marks	4	0
		Suggesting improvements	3 marks	2	0

MMO = Manipulation, measurement and observation

Collection = Successful collection of data and observations

Decisions = Decisions relating to measurements or observations

PDO = Presentation of data and observations

Recording = Recording data and observations

Display = Display of calculation and reasoning

Layout = Data layout

ACE = Analysis, conclusions and evaluation

Interpretation = Interpretation of data or observations and identifying sources of error

Conclusions = Drawing conclusions

Improvements = Suggesting Improvements

Page 3	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

Qι	Question		Sections	Learning outcomes	Indicative material	mark
1	(a)		MMO Decisions  ACE Conclusions	<ul> <li>Decide how many tests, measurements or observations to perform</li> <li>Make and record sufficient, accurate measurements and observations</li> <li>Draw conclusions from interpretations of observations, data and calculated values</li> </ul>	2 very simple tests ethanol emulsion, shake = clear AND benedicts + heat = red/ orange/yellow (R green); reducing sugar (R glucose/ other unqualified sugar);	1
	(b)	(i)	MMO Decisions	<ul> <li>decide how many tests, measurements or observations to perform</li> <li>make measurements or observations that span the largest possible range within the limits either of the equipment provided or of the instructions given</li> <li>make quantitative measurements or qualitative observations that are appropriately distributed within this range</li> </ul>	for room temperature: at least 2 and not more than 4 readings, each of at least 10 seconds and nor more than 60 seconds;	1

Page 4	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

	(ii)	MMO collection	set up apparatus correctly     follow instructions given in	data reported as bubbles per unit time for at least two	1
		MMO	the form of written instructions or diagrams • decide how many tests,	temperatures; decide to investigate three or	1
		decisions	measurements or observations to perform • replicate readings or	more temperatures and to replicate readings	
		MMO collection	<ul> <li>observations as necessary</li> <li>make and record sufficient, accurate measurements</li> </ul>	at least three temperatures investigated, and at least two	1
		PDO recording	<ul> <li>and observations</li> <li>present numerical data,</li> <li>values or observations in a single table of results</li> </ul>	replicate readings made; all data recorded in a single table with appropriate means to record bubbling rate per	
			draw up the table before taking readings/making observations, so that	unit time, replicated, at more than one temperature; column headings that include	2
			candidates can record directly into the table, to avoid the need to copy up	quantities and unit where appropriate (such as temperature/°C, number of	
			their results  Include in the table of results, if necessary, columns for raw data, for	bubbles in 10 seconds);	
			calculated values and for deductions  use column headings that		
			include the quantity and the unit (as appropriate) and that conform to accepted		
		PDO layout	scientific conventions     choose a suitable and clear method of presenting the	most data recorded in a table;	
			data, e.g. tabulations, chart, graph, drawing or mixture of methods of presentation		1

Page 5	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

(c)	(i)	ACE interpretation	evaluate the effectiveness of control of variables and thus the confidence with which conclusions might be drawn	no attempt made to control it so not well controlled /distilled water used, so no acid or alkali added, but not well controlled/no buffer added so not well controlled/yeast contains proteins/buffers/weak acids + salts that might help buffer the solution a little;	1
	(ii)	ACE interpretation	identify the most significant sources of error in an experiment	Two from: IDEA OF bubbles might vary in size/ temperature change might cause gas inside tube to change volume/one example of limited accuracy of measuring equipment e.g. syringe/AVP;	1
(d)	(i)	MMO Decisions	replicate readings or observations as necessary (individual readings or observations should be repeated where they appear to be anomalous)	something has gone wrong with the apparatus / the gas bubbles have leaked out somewhere / AVP (accept reading anomalous / not reliable unqualified);	1
	(ii)	PDO display	<ul> <li>show their working in calculations, and the key steps in their reasoning</li> <li>use the correct number of significant figures for calculated quantities</li> </ul>	4.9 with appropriate working shown; R no working shown R more than two significant figures	1

Page 6	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

	(iii)	PDO layout	select which variable(s) to plot and plot appropriately on clearly labelled x- and y-axes     plot all points or bars to an appropriate accuracy     follow the IOB recommendations for putting lines on graphs	independent variable (temperature) on x-axis, dependent variable (mean number of bubbles min <sup>-1</sup> ) on y-axis AND axis labels appropriate (accept ecf from table if already penalised in 1 (b) (ii)); scale should be chosen so that data spans at least half of the width and height of the grid AND scale appropriate such as 1:10, 1:5 or 1:2 (R awkward scales such as 3:10, 7:10, 8:10) (scale does not need to start at 0); data plotted accurately to within 1 mm, using crosses or circle-with-dot AND points joined with straight ruled lines OR fine curve drawn through the data points, not extrapolated beyond the first or last point;	3
(e)		ACE Conclusions	•draw conclusions from an experiment, giving an outline description of the main features of the data, considering whether experimental data supports a given hypothesis, and making further predictions	at low temperatures an increase in temperature increases bubbling rate, AND at high temperatures an increase in temperature decreases bubbling rate/AW;	1

Page 7	Page 7 Mark Scheme		Paper
	GCE A/AS LEVEL – 2007	9700	31

(f)	ACE Conclusions	draw conclusions from an experiment, giving an outline description of the main features of the data, considering whether experimental data supports a given hypothesis, and making further predictions	IDEA OF at low temperatures the data supports the student's hypothesis AND above 48 °C/at high temperatures the hypothesis is not supported/the rate drops as temperature increases; prediction including student's hypothesis for low temperatures PLUS at high temperatures, as temperature increases, the rate of production of gas will decrease/AW;	2
(g)	ACE Improvements	suggest modifications to an experimental arrangement that will improve the accuracy of the experiment or the accuracy of the observations that can be made, including the use of new methods or strategies to investigate the question describe such modifications clearly in words or diagrams	accept improvements that would enhance the reliability or accuracy of the experiment – three in outline or one or two explained – could be related to errors identified earlier or others collect gas; measure its volume accurately; e.g. of specific method of doing so such as inverted burette over water/gas syringe; use more replicates/repeat more times at each temperature; use more temperatures/ specified wider range between 0 and 100 °C; use more accurate measuring devices/one named specific measuring device; use a buffer to control pH/ other specific means to control a plausible variable; AVP;;	2

Page 8	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

2	(a)	(i)	MMO Collection	set up apparatus correctly     use their apparatus to     collect an appropriate     quantity of data or     observations, including     subtle differences in     colour or other properties     of materials	Rancunculus root t.s. recognisable in drawing (large circle containing smaller circle containing star- shaped region); proportions of stele/root diameter acceptable (between 1:5 and 1:10) AND at least 4 tissues shown (epidermis, parenchyma, endodermis, xylem, phloem);	2
		(ii)	MMO Collection	<ul> <li>make measurements         using millimetre scales,         graticules, protractors,         stopwatches, balances,         measuring cylinders,         syringes, thermometers,         and other common         laboratory apparatus.</li> <li>show their working in         calculations, and the key         steps in their reasoning</li> </ul>	correct measurement of line shown on drawing to within 1 mm AND measurement of diameter of specimen between 1.5 and 4 mm, to no more than 0.5 mm reported accuracy; working shows measurement from drawing divided by measurement from slide;	1
		(iii)	ACE Interpretation	<ul> <li>estimate, quantitatively, the uncertainty in quantitative measurements</li> <li>express such uncertainty in a measurement as an actual or percentage error</li> </ul>	their reported measurement $\pm$ 0.5 mm (accept answers between $\pm$ 0.2 mm and $\pm$ 1.0 mm)	1
		(iv)	ACE Interpretation	show an understanding of the distinction between systematic errors and random errors	ruler made with incorrect intervals/user not viewing at right angles/AVP;	1

Page 9	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

(b)		MMO Collection  MMO Decisions  PDO Layout	<ul> <li>use their apparatus to collect an appropriate quantity of data or observations, including subtle differences in colour or other properties of materials</li> <li>decide how many tests, measurements or observations to perform</li> <li>make measurements or observations that span the largest possible range within the limits either of the equipment provided or of the instructions given</li> <li>make quantitative measurements or qualitative observations that are appropriately distributed within this range</li> <li>choose a suitable and clear method of presenting the data, e.g. tabulations, chart, graph,</li> </ul>	at least half of area of available space used to represent/describe a number of cells; drawings/descriptions of cells including starch granules, cell walls and air spaces between corners of the cells; at least three and no more than six cells drawn/ described; largest cell drawn/described at least twice the size of smallest; cells with a range from 2 or less up to 10 or more starch grains; including both cells with air spaces between the corners and those without;  drawing used to represent observations – clear outline drawings, sharp pencil and no shading;	2 max 3
			drawing or mixture of methods of presentation	no snaung,	
(c)	(i)	PDO layout	<ul> <li>choose a suitable and clear method of presenting the data, e.g. tabulations, chart, graph, drawing or mixture of methods of presentation</li> </ul>	table used to present data; (R comparative lists without lines to divide information)	1

Page 10	Mark Scheme	Syllabus	Paper
	GCE A/AS LEVEL – 2007	9700	31

	(ii)	MMO Collection  PDO Recording	<ul> <li>use their apparatus to collect an appropriate quantity of data or observations, including subtle differences in colour or other properties of materials</li> <li>present numerical data, values or observations in a single table of results</li> <li>draw up the table before taking readings/making observations, so that candidates can record directly into the table, to avoid the need to copy up their results</li> <li>record raw readings of a quantity to the same degree of precision and observations to the same level of detail</li> </ul>	Give at least 4 comparisons, including at least one similarity and at least one difference, and including one subtle judgement (judgement involving more than just size, colour or shape); all observations and comparisons recorded in a single table with difference(s) recorded to the same level of precision (e.g. sizes recorded in mm) or detail (e.g. stele 40% of total width of S3 vs. stele 8% of total width of specimen S4);	1
	(iii)	ACE Interpretation	<ul> <li>describe and summarise the key points of a set of observations</li> </ul>	central stele/named feature (e.g. xylem/tubular cells);	1
(d)		MMO Decisions ACE Interpretation	<ul> <li>make and record sufficient, accurate measurements and observations</li> <li>describe and summarise the key points of a set of observations</li> </ul>	correctly label xylem on both pictures;  pick out at least one valid reason for each decision (e.g. Fig. 2.1 thick cell walls, Fig.2.2 end walls of cells absent);	1

Centre Number	Candidate Number	Name

# UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Level

BIOLOGY 9700/04

Paper 4 Structured Questions

For Examination from 2007

Specimen Paper

1 hour 15 minutes

Additional Materials: Answer Booklet/Paper

### **READ THESE INSTRUCTIONS FIRST**

Write your Centre number, candidate number and name on all the work you hand in. Write in dark blue or black pen.

Do not use staples, paper clips, highlighters, glue or correction fluid.

#### **Section A**

Answer all questions.

#### **Section B**

Answer one question.

Write your answer on the separate Answer Booklet/Paper.

At the end of the examination, fasten all your work securely together.

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

[2]

	, ,	0				/D _ \
1	(a)	State what is meant by	/ the term	respiratory	/ quotient (	(KQ)

[1]

**(b) (i)** Complete the following equation for the aerobic respiration of the respiratory substrate A.

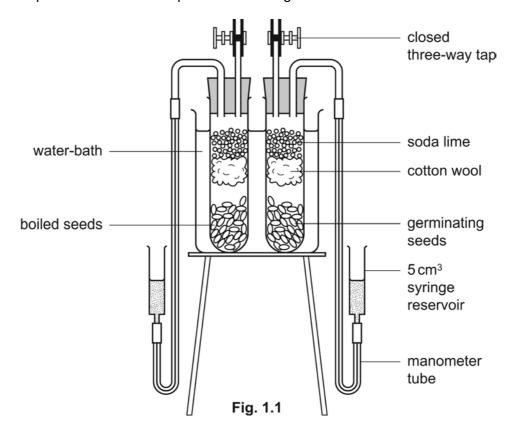
$$C_{18} H_{36} O_2 + 26 O_2$$
 + [2]

(ii) Calculate the respiratory quotient (RQ) of this respiratory substrate.

(c) Explain the significance of the different values that may be obtained of RQ.

[2]

Two respirometers were set up as shown in Fig. 1.1.



(d) Outline how this apparatus is used to measure the rate of oxygen uptake mass of germinating seeds.	e by a known
	[4]
(e) Explain how the apparatus could be modified to measure the RQ of seeds.	the germinating
	[2]
(f) Explain why an increase in temperature from 15 °C to 25 °C will increase oxygen uptake in germinating seeds.	ease the rate of
	[2]
	[Total: 15]

[Total: 8]

2 Fig. 2.1 shows the main stages of the Calvin cycle.

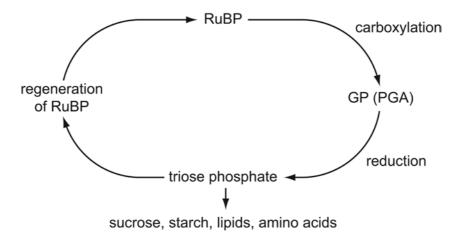


Fig. 2.1

(a)	State precisely where the Calvin cycle occurs in plant cells.	[1]
(b)	Describe how carbon dioxide is fixed in the Calvin cycle.	[,]
(c)	Explain how the products of photophosphorylation are used in the Calvin cycle.	[2]
		[3]
(d)	Explain what initially happens to the concentration of RuBP and GP if the supply of carbon dioxide is reduced.	
	RuBP	
	GP	
		[2]

3 Scallops, which are bivalve molluscs, are important commercially throughout the world. The marine bay scallop, *Agropecten irradians*, has three distinct shell colours, yellow, orange and black. The shell colour is controlled by a gene with three alleles, yellow, **S**<sup>y</sup>, orange, **S**°, and black, **S**<sup>b</sup>.

Scallops are hermaphrodite and are able to fertilise themselves to produce offspring.

Single mature adult specimens of yellow, orange and black scallops were collected and kept in separate tanks of seawater until they produced young. The young were then scored for shell colour. The results were as follows:

yellow scallop – 25 yellow and 8 black orange scallop – 31 orange and 9 black black scallop – 27 black

(a)	Explain the results from the orange and black scallops, using the symbols given.
	[6]
(b)	Orange scallops are more valued for human consumption.
	Describe how a marine biologist could produce a pure-breeding line of orange scallops for commercial exploitation using the offspring from the single orange scallop.
	[2]
	Total: 81

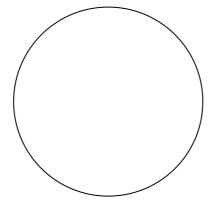
4 (a) Fig. 4.1. shows the above-ground parts of the orchid *Masdevallia caudata*.



Fig. 4.1

(i)	State <b>one</b> feature, visible in Fig. 4.1, that indicates that this plant is angiospermophyte.	an
		[1]
(ii)	State <b>two</b> features, visible in Fig. 4.1, that indicate that this plant is monocotyledon.	а
	1	
	2	[2]
(iii)	Describe the type of root system that you would expect to be present on this pla	nt.
		[2]

(iv) On the outline below of a transverse section of the stem of this orchid, sketch the positions of the vascular bundles that you would expect to find.



[1]

**(b)** Table 4.1 lists three plant phyla and three features that may be found in them.

Complete the table by placing a tick to indicate when a feature is present in a phylum and a cross to indicate when it is not. (Do not leave any boxes blank).

Table 4.1

feature	bryophytes	filicinophytes	coniferophytes
dominant stage is diploid sporophyte			
vascular tissue present			
xylem vessels present			

[5]

Describe two ways in which the life cycle of an angiospermophyte complete adaptation to life on dry land than that of a bryophyte.	shows	more
1		
		••••••
2		
		[4]
	[Tota	al: 15]
	complete adaptation to life on dry land than that of a bryophyte.  1	complete adaptation to life on dry land than that of a bryophyte.  1  2

5 Fig. 5.1 outlines the way in which the gene for human insulin is incorporated into plasmid DNA and inserted into a bacterium.

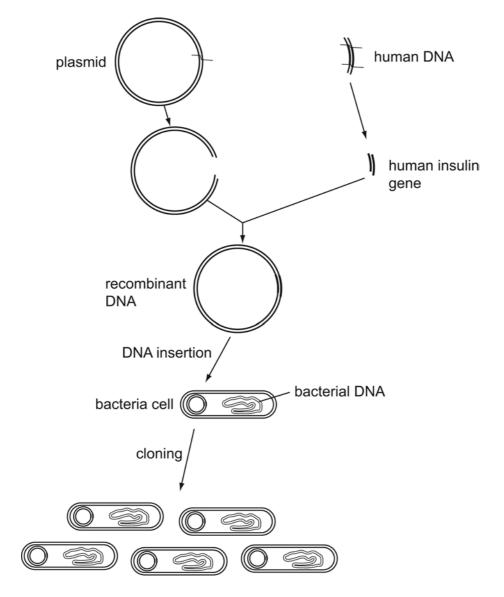


Fig. 5.1

(a)	Describe how the plasmid DNA is cut.	
		•
	[3]	
(b)	Explain how the human insulin gene is joined to the plasmid DNA.	
		•
		•
	[3]	
(c)	List <b>two</b> advantages of treating diabetics with human insulin produced by genetic engineering.	;
	1	•••
		•••
	2	
		2]
	[Total: 8]	

6	(a)	Describe the roles of barley and yeast enzymes in beer production.					
							[3]
	(b)	Mo				ers of low energy conte	
	(6)	mo	re popula	r. Light beers	have a low starch case after the mashing	ontent. This is achieved	d by the addition
		(i)	Explain t	he advantage	of using immobilised	d enzymes in this proces	SS.
							[0]
		/::\		at of two differ	out toward of immarkill	in a definition of a second se	the hydrolysis of
		(11)				ised fungal amylase on tions, starch is not a lim	
					Table 6.1		
			time/b	mass of malt	ose produced/g		
			time/h	ume/m	α amylase	β amylase	
				0	0	0	
				1	0.05	0.05	
				2	0.20	0.10	
				3	0.60	0.20	
					le 6.1, explain which rs with a low starch c	of these enzymes woul	d be used in the
							[2]
							[Total: 8]

7	(a)	Describe the structural features of wind pollinated plants such as grasses.
		[4]
	(b)	State two advantages of self pollination and two advantages of cross pollination.
		self pollination
		1
		2
		cross pollination
		1
		2
		[4]
		[Total: 8]

8 (a) Name the precise sites of production in the human male of the following hormones:

(ii) luteinising hormone (LH) or interstital cell stimulating hormone (ICSH);

(i) follicle stimulating hormone (FSH);

(iii) testosterone.

[3]

**(b)** Fig. 8.1 shows the concentration of the hormones FSH, LH (ICSH) and testosterone in the blood of a human male at different ages.

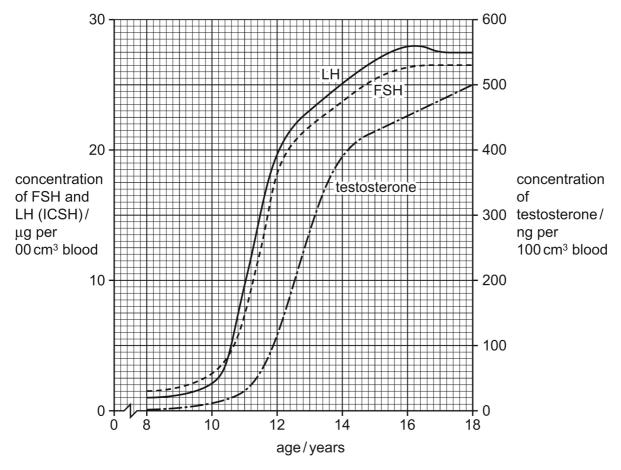


Fig. 8.1

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dable 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g  10 2.0  12 3.0  14 8.0  16 18.0  18 28.0  //tth reference to Table 8.1, calculate, showing your working in each case,	(i)	-	and explain the changes in concentration of	
able 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g 10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;				•••••
able 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g 10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;				
able 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g 10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;				
able 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g 10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;				
able 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g 10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;				•••••
able 8.1 shows the mean mass of the human testis at different ages.  Table 8.1  age/years mean mass of human testis/g 10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;				••••
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;	(ii)	testosterone.		
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;				
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;				
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;				
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;				
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;				
Table 8.1    age/years   mean mass of human testis/g   10   2.0     12   3.0     14   8.0     16   18.0     18   28.0     28.0    /ith reference to Table 8.1, calculate, showing your working in each case,   the absolute growth rate of the testis between ages 14 and 18 years;	·······	0.4 above the many many of t	he he was to still at different area	•••••
age/years mean mass of human testis/g  10 2.0  12 3.0  14 8.0  16 18.0  18 28.0  With reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;	able			
10 2.0 12 3.0 14 8.0 16 18.0 18 28.0  With reference to Table 8.1, calculate, showing your working in each case, 14 the absolute growth rate of the testis between ages 14 and 18 years;		Ta	able 8.1	
12 3.0 14 8.0 16 18.0 18 28.0  //ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;		age/years	mean mass of human testis/g	
14 8.0 16 18.0 28.0  /ith reference to Table 8.1, calculate, showing your working in each case, the absolute growth rate of the testis between ages 14 and 18 years;		10	2.0	
16 18.0 28.0  With reference to Table 8.1, calculate, showing your working in each case,  the absolute growth rate of the testis between ages 14 and 18 years;		12	3.0	
/ith reference to Table 8.1, calculate, showing your working in each case,  the absolute growth rate of the testis between ages 14 and 18 years;		14	8.0	
/ith reference to Table 8.1, calculate, showing your working in each case,  the absolute growth rate of the testis between ages 14 and 18 years;		16	18.0	
the <b>absolute</b> growth rate of the testis between ages 14 and 18 years;		18	28.0	
the <b>absolute</b> growth rate of the testis between ages 14 and 18 years;				
	Vith :	eference to Table 8.1, calculat	e, showing your working in each case,	
	i) tł	ne <b>absolute</b> growth rate of the	testis between ages 14 and 18 years:	
	.,	io abbolato growal rato or the	teetie between ages 11 and 10 years,	
	1.			
	181			
i) the <b>relative</b> growth rate of the testis between ages 14 and 18 years.				•••
		ne <b>relative</b> growth rate of the te	estis between ages 14 and 18 years.	
	 ii) tl			
	 ii) tl			
	 ii) tl			

# Section B Answer one question

9 (a) Explain how a synapse functions. [9]
(b) Describe the role of glucagon in regulating blood glucose. [6]
10 (a) Describe why variation is important in natural selection. [6]
(b) Explain the role of isolating mechanisms in the evolution of new species. [9]
[7otal: 15]

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#### **UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS**

Specimen for 2007

## **GCE A LEVEL**

# MARK SCHEME

**MAXIMUM MARK: 100** 

SYLLABUS/COMPONENT: 9700/04

BIOLOGY STRUCTURED QUESTIONS



		Page 2	Mark So		Syllabus	Paper
			GCE A LEV	/EL – 2007	9700	4
1	(a)		of carbon dioxide given off; of oxygen taken up	R amount	A moles	[1]
	(b)	(i) 18H <sup>2</sup> O; 18CO <sup>2;</sup>				[2]
		(ii) 18/26; = 0.69 –	0.70;	allow 2 marks fo	or correct ar	nswer [2]
	(c)	carbohydrate	substrate; 1 some anaerobic respiration e 1/protein 0.9/fat 0.7 ;; metabolic processes using ox	2 out of 3		[2 max]
	(d)	record level of change in known repeat; open clip and ref. units; ref. to boiled	seeds as a control; absorbs carbon dioxide giver	n off;		[4 max]
	(e)		iment/ref. to comparison; er manometer rose or fell;			[2 max]
	(f)	ref. effect of t	temperature on <u>enzymes in re</u> ffect of temperature e.g. incre		gy/more sub	
			th activation energy;			[2 max]
		101. 10 & 10	-			Total: 15
2	(a)	stroma of chl	oroplast;			[1]
	(b)		th (5C compound) RuBP; able 6C compound/forms 2 mo rubisco;	olecules of (3C) GP;		[2 max]
	(c)	(reduced NA) ref. use of A7	OP and ATP; rce of energy; DP is for) reduction of GP(PG TP in regeneration of RuBP; e of phosphate/phosphorylatio	, , , ,	);	[3 max]
	(d)	due to reduce	nulates/goes up; ed combination with CO <sub>2</sub> /AW; wn/not as much being formed		or GP, not b	oth
		due to conve		,		[2 max]
						Total: 8

	Page	3			Mark S	Scheme			Syllabus	Paper
					GCE A LE	VEL – 200	7		9700	4
(a)	Eitl If g	<u>ner</u> enetic dia	agram u	sed	Pena	alise once	e for inco	rrect syr	nbols	
					oran	ge domina	ant to blac	k (for coı	nverse);	
ora	nge s	scallop								
-	ents netes	3	S°	S°Sb	S <sup>b</sup>	X	S°	S°Sb	S <sup>b</sup>	
ger	notyp	е	S° S°		S° Sb		S° Sb		S <sup>b</sup> S <sup>b</sup>	
phe	enoty	ре			orange				black	
bla	ck sc	allop		$S^b S^b$		Χ		$S^b S^b$		
par	ent									
gar	netes	3			(	S <sup>b</sup>		$S^b$	)	
ger	notyp	е				S <sup>b</sup> S <sup>b</sup>				
phe	enoty	ре				black				
	Or If to	ext explai	nation g	iven						
	ora (be- link blac	nge dominge are he cause) referenced to refe	eterozyg <sup>f</sup> . 3:1 rati atio; nozygou	ous; o; s;	·					1
(b)	son	ne will pro	duce onl	y orange				orange	with black;	[2ma
										Total
(a)	(i)	flowers								
	(ii)	<u>3</u> -petalle parallel v					ignore	elongate	ed leaves	
	(iii)	adventition		:						

no tap root;

(iv) bundles scattered and not in a ring;

[2 max]

[1]

Page 4	Mark Scheme S		Paper
	GCE A LEVEL – 2007	9700	4

	Bryophyta	Filicinophyta	Coniferophyta
dominant stage is diploid sporophyte	х	✓	✓
vascular tissue present	х	✓	✓
xylem vessels present	Х	х	х

half mark per correct box, round up

(b)

[5]

(c) assume statement refers to angiiospermophyte unless otherwise stated.

gametophytes/male gametes, inside pollen grain; protected from desiccation/can be dispersed over wide area;

internal fertilisation/fertilisation described; not dependent on water/male gametes do not swim;

young sporophyte/embryo, develops within seed; not dependent on gametophyte/can lie dormant for long periods/can survive dry conditions

[4 max]

Total: 15

5 (a) restriction (endonuclease) enzyme;

named example; e.g. EcoR1 specific, sequence of bases/point; ref. to sticky ends/exposed bases;

[3 max]

(b) sticky ends added to insulin gene;

ref. to complimentary base pairing/C and G bases pair up;

ref. H bonds;

(DNA) ligase;

formation of phosphodiester bond/seals sugar phosphate backbone;

[3 max]

(c) identical to human insulin (ref. to bovine/porcine insulin used previously);

ref. to reduced immune response/side effects;

cheaper to produce;

more rapid response;

pure/uncontaminated;

regular production not dependent on livestock;

ethical issues:

AVP; e.g. tolerance

[2 max]

Total: 8

6 (a) amylase;

hydrolyses starch;

to maltose;

ref. α and β amylase;

maltose to glucose;

maltase;

anaerobic breakdown/glycolysis, of sugar;

into ethanol and carbon dioxide;

[3 max]

Page 5		Mark Scheme	Syllabus	Paper
		GCE A LEVEL – 2007	9700	4
(b)	enzyme enzyme idea of AVP; e.	ase; naltose produce;		[3 max] [2 max] Total: 8
' <b>(</b> a)	flowers held	oduced; a; gma;		[4 max]
(b)	effective in he.g. high mo	dely scattered; harsh environments; ountains max 2 ation iation;		[4 max]
				Total: 8
3 (a)	(ii) anterior	r_pituitary gland; r_pituitary gland; tial cells/Leydig cells, (of testis); y + pituitary + testis = 1)		[3]
(b)	plateau ref. figu triggere from hy steep ri LH/ICSI	ant rise in both at age 10-12 years; u in both at ages 16/17 years; ures; ed by GnRF; ypothalamus; ise triggers puberty; sH stimulates synthesis of testosterone; imulates spermatogenesis;		[4 max]

Page 6	Mark Scheme S		Paper
	GCE A LEVEL – 2007	9700	4

(ii) significant rise at age 10/11-14 years;

rise less steep age 14-18 years;

ref. figures;

triggered by rise in LH(ICSH)

testis increases in size at the same time;

responsible for secondary sexual characteristics;

[4 max]

(c) (i) 
$$\frac{28.0 - 8.0 \, g}{4y}$$
 = 5; g per year; [2]

(ii) 
$$\frac{20}{4} \times \frac{1}{8}$$
 or  $\frac{5}{8}$ ; = 0.625 (0.63 s.f.); [2]

Total: 15

9 (a) Explain how a synapse functions.

[9]

(b) Describe the role of glucagon in regulating blood glucose.

[6]

- (a) 1 depolarisation/action potential;
  - 2 of presynaptic membrane/synaptic knob;
  - 3 opening calcium ion channels;
  - 4 calcium ions in;
  - 5 vesicles containing transmitter/acetylcholine;
  - 6 fuse with membrane;
  - 7 contents emptied into synaptic cleft/exocytosis;
  - 8 transmitter/acetylcholine diffuses across synaptic cleft;
  - 9 transmitter/acetychloine binds to receptor; **R** protein channel
  - 10 on post synaptic membrane;
  - 11 Na<sup>+</sup> channels open/NA<sup>+</sup> enters;
  - 12 depolarises post synaptic membrane;
  - 13 action potential set up/impulse transmitted
  - 14 breakdown/hydrolysis of transmitter/acetylcholine by enzyme/cholinesterase; [9 max]
- (b) 15 when blood glucose levels low;
  - 16 glucagon released from alpha cells (in pancreas);
  - 17 (acts on ) liver (cells);
  - 18 breakdown of glycogen to glucose;
  - 19 use of fatty acides in respiration;
  - 20 production of glucose from other compounds/fats/amino acids/gluconeogenesis;
  - 21 liver releases glucose into blood;
  - 22 alucose levels rise/return to normal:
  - 23 switching off glucagon secretion;
  - 24 antagonistic to insulin;

[6 max]

Total: 15

Page 7	Mark Scheme	Syllabus	Paper
	GCE A LEVEL – 2007	9700	4

- 10 (a) 1 ref. continuous/discontinuous variation;
  - 2 genetic/inherited variation;
  - 3 variation in phenotype/characteristics/AW;
  - 4 (can be due to) interaction of genotype and environment;
  - 5 e.g. of characteristic that influences survival;
  - 6 ref. intraspecific competition/struggle for existence;
  - 7 those with favourable characteristics survive/AW;
  - 8 pass on favourable characteristics to offspring;
  - 9 those with disadvantageous characteristics die;

[6 max]

- (b) 10 ref. to definition of species;
  - 11 ref. allopatric;
  - 12 geographical isolation;
  - 13 ref. to examples e.g. islands/lakes/mountain chains/idea of barrier;
  - 14 ref. to example organism;
  - 15 ref. to populations prevented from interbreeding;
  - 16 isolated populations subjected to different selection pressures/conditions;
  - 17 over time sufficient differences to prevent interbreeding;
  - 18 ref. sympatric;
  - 19 ref. to reproductive isolation;
  - 20 ref. behavioural barriers (within a population);
  - 21 e.g. day active/night active;
  - 22 correct ref. to gene pool;
  - 23 change to allele frequencies;

[9 max]

Total: 15

Centre Number	Candidate Number	Name

# UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Subsidiary Level and Advanced Level

BIOLOGY 9700/05

Paper 5 Planning, Analysis and Evaluation

For Examination from 2007

Specimen Paper

Candidates answer on the Question Paper. No Additional Materials are required. 1 hour 15 minutes

#### **READ THESE INSTRUCTIONS FIRST**

Write your Centre number, candidate number and name on all the work you hand in. Write in dark blue or black pen.

Do not use staples, paper clips, highlighters, glue or correction fluid.

Answer all questions.

At the end of the examination, fasten all your work securely together.

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

1 (a) The rate of respiration in two tissues, **A** and **B** was measured using DCPIP as an indicator. 50 g of each tissue was ground into paste using 10 cm<sup>3</sup> of ice cold buffer and made into a suspension with 250 cm<sup>3</sup> of buffer solution. The two suspensions were placed into a water bath at 20 °C.

Two sets of ten tubes, each containing 10 cm³ of buffer and 1 cm³ of DCPIP, were placed into separate water baths at 20 °C and left for 10 minutes.

To one set of ten tubes, 0.5 cm³ of suspension **A** was added. To the other set of ten tubes, 0.5 cm³ of suspension B was added. The time taken for DCPIP to become colourless was measured separately in each tube.

(i)	State the dependent and independent variable in this investigation.
	independent variable
	dependent variable [1]
(ii)	Identify the key variables that have been controlled in this investigation.
	For each variable, describe how it has been controlled.
	[3]
(iii)	Suggest how the colourless end point of the DCPIP might have been standardised.
	[1]

Table 1.1

	time taken for DCPIP to become colourless/s												
1 2 3 4 5 6 7 8 9 10									10				
Tissue A	55	56	59	54	52	56	55	55	59	59			
Tissue B	Tissue B 125 126 122 126 122 119 121 123 124 125												

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**(b)** The results of this investigation are shown in Table 1.1.

(i)	U	se	t t	ne	fc	orr	ทเ	ıla	ae	b	el	ΟV	v 1	to	Ca	alc	cul	la	te	th	ıe	si	taı	าด	laı	rd	е	rr	or	fo	or	е	ac	h	of	f t	he	e ti	iss	sue	es	te	est	te	d.
	S	taı	nd	ar	ď	de	ev	ia	tio	n	(s	s)								Standard error S <sub>M</sub>																									
	s	=	$\sqrt{2}$	Σ(	x n		<u>x</u> ) 1	)2	-																						3	S <sub>M</sub>	, =	= -	S √r	= 7									
	;	Sta	an	da	arc	d∈	err	OI	r fo	or	Si	an	np	ol∈	<b>.</b> A	1				•••				•••		•••	••••				•••	•••	••••				••••			••••			•••	•••	,
	,	Sta	an	da	arc	d e	err	OI	r fo	or	S	an	np	ole	) E	3					••••				•••		•••	•••					••••										[	4]	
(ii)		se						tc	þ	olo	ot	а	b	a	r	ch	ar	t	of	tl	he	1	ne	ea	n	re	es	ul	ts	6	an	ıd	S	ta	nc	la	rd	e	rr	or	fc	or	th	iis	,
					Ŧ			-					-					-								-		1		1				+				H					-		F
			H		Ŧ														H				Ŧ							1				1			#	Ħ	Ŧ	1	Ħ	Ħ	+		Ŧ
	Ħ		Ħ		ŧ		H		Ħ						Ξ		Ħ	ŧ					Ξ				Ħ			1								Ħ							Ė
				H	+		H	+	Ħ	Ė					Ŧ			ŧ		Ė			ŧ					+		+				ŧ								$\exists$	1		Ė
	H		H	H	#			+	H				+		Ŧ		H	+	H							+		+		+				+		H	#	Ħ				$\parallel$	+		F
	H		H	H	Ŧ					1								+	H	F	+		Ŧ			-				1				Ŧ			=	Ħ	Ŧ				+		F
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#### **2** Fig. 2.1 shows the structure of a root tip.

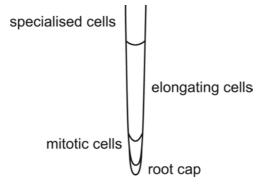


Fig. 2.1

Radicles of seedlings constantly produce new cells by mitosis. As these cells develop they become elongated by absorbing water and the vacuole expanding and causing the cell wall to stretch. Growth regulators control the plasticity of the cell wall so it is able to stretch.

The developing radicles of seedlings placed horizontally respond by curving and growing in the direction of gravity.

One hypothesis to explain this curvature is that the root cap contains gravity receptors that causes changes in the distribution of auxin secreted by the root tip.

Describe how this hypothesis could be tested.
[10]

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**3** Fig. 3.1 shows genetic fingerprints made from DNA samples of a number of different mammals.

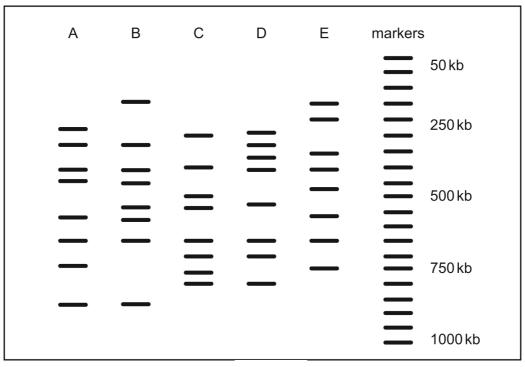


Fig. 3.1

Explain how the genetic finger print provides evidence that these mammals share a common ancestor.
 [2]
 Use the information in the diagram to explain the evolutionary relationship between these mammals.
 [5]

[Total 7]

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#### **UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS**

Specimen for 2007

## **GCE A LEVEL**

# MARK SCHEME

**MAXIMUM MARK: 30** 

SYLLABUS/COMPONENT: 9700/05

BIOLOGY PRACTICAL



Page 2	Mark Scheme	Syllabus	Paper
	GCE A LEVEL – 2007	9700	5

Skill	Total	Breakdown of mark	<b>〈</b>	Question	Question	Question
	marks	expectations	expectations			3
Planning	15	<b>PP</b> Defining the	5	5		
	marks	problem				
		PM Methods	10		10	
Analysis,	15	<b>DD</b> Dealing with	8	8		
conclusions	marks	data				
and		<b>E</b> Evaluation	4			4
evaluation		C Conclusion	3			3

The abbreviations in bold are used to identify the marks in the mark scheme below

1 (a) (i) tissue source,

time (for DCPIP to decolourise);

1 PP

(ii) temperature – water bath at 20 °C

pH - buffer solution

osmotic balance - buffer solution

relative concentrations indicator, tissue sample – standard volumes of each quantity of tissue sample – same mass and volumes buffer used

1-2 decognised 1
3-4 recgnised 2
5 recognised 3

(iii) 10 cm<sup>3</sup> buffer,

1 cm<sup>3</sup> water/buffer,

0.5cm<sup>3</sup> tissue sample (same as that being tested)

1 PP

3 PP

**(b) (i)**  $s= 2.23, S_M = .1.58;$ 

$$s= 2.49, S_M = 0.79;$$

4 DD

(ii) axes correct orientation;

both plots correct (means 55.9 sand 122 s)

both standard error bars correctly plotted;

3 DD

(iii) results for sample B more accurate than sample A as there is less spread

1 DD

Total 13 PP and DD

Page 3	Mark Scheme	Syllabus	Paper
	GCE A LEVEL – 2007	9700	5

#### 2 Independent variables

radicles all same approximate age/ length/measured;

half left with root caps intact/half root caps removed;

one set with root caps and one set without root caps placed horizontally;

one set with root caps and one set without root caps left vertically;

suitable number of seedlings – batches not less than 25;

max 2

Dependent variable

suitable method of measuring – accuracy using mm/callipers

before and after standard time (24-48 hours;)

method of measuring curvature;

max 2

Key variable and controls:

ref. to standard conditions/examples of conditions for germination(any2);

radicles all same approximate age/ length/measured;

ref. to standard conditions and explantion re. growth of radicles;

indentification of the 3 sets of controls and their role;

max 2

Apparatus and quantities;

suitable method of germinating seedlings – petridishes+ paper/cotton wool;

germination conditions standardised;

suitable number of seedlings – batches not less than 25;

Risks and precautions

ref. to safety - cutting and care needed;

max 2

Data presentation

table showing results presentation –suitable headings with units;

Use of data

ref. to how results might confirm/disprove hypothesis;

(idea that root cap removed – no response but growth continues

root cap intact responds if horizontal by uneven growth);

max 2

10 PM

Page 4	Mark Scheme	Syllabus	Paper
	GCE A LEVEL – 2007	9700	5

3 (a) DNA fragments of similar size/ positions for all types of mammal;

ref. to bands at 400kband 700kb;

2 E

**(b)** A and B more closely related to each other 3 bands in common

around 300 kb, 55-6000 kb, 800 kb

have separated most recently in evolution

E is distantly related – has only one other bandt/350 kb in common with D;

and no others with A and B;

separated early in evolution from any of the other types of mammal

C and D closely related each other as 2 bands in common;

around 700 kb and 800 kb;

A and B/C and D more distantly related/separated in evolution;

D has common band/ 300-350kb with A and B

C has common band/500-550kb with B and D

5 EC

Total 7 EC