

UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS General Certificate of Education Advanced Level

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		

BIOLOGY 9700/53

Paper 5 Planning, Analysis and Evaluation

May/June 2013

1 hour 15 minutes

Candidates answer on the Question Paper.

No Additional Materials are required.

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 ink.

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

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

DO NOT WRITE IN ANY BARCODES.

Answer all questions.

Electronic calculators may be used.

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.

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1			
2			
Total			

This document consists of 7 printed pages and 1 blank page.



1 (a) A group of students was given the task of planning a method to find the effects of the growth regulator gibberellin (GA) on the germination of maize grains.

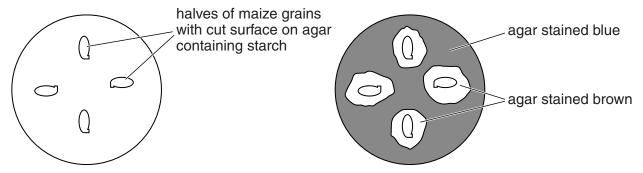
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The students looked up information about investigations into the germination of maize. The information that the students found is listed below.

- Maize needs to be soaked in water for 24 hours to stimulate germination.
- Maize starts to germinate 48–72 hours after soaking.
- GA promotes germination by activating a gene that causes the synthesis of the enzyme amylase.
- The range of concentrations at which GA is found in plants is between $1-10\,\mu\text{mol}\,\text{dm}^{-3}$.
- Amylase is released by the aleurone layer into the endosperm of the maize grain to hydrolyse the starch reserves.
- The activity of amylase can be estimated by cutting the maize grains lengthways into two halves and placing the cut sides onto agar containing starch in Petri dishes.
- After incubation at a constant temperature, iodine solution is used to test the agar for the presence of starch.

The students carried out a preliminary investigation using maize grains that had been soaked in a 3 mmol dm⁻³ solution of GA.

Fig. 1.1 shows the arrangement of cut maize grains that the students decided to use for their main investigation and the results of testing the agar for starch using iodine solution after incubation.



arrangement of maize grains on agar containing starch in a Petri dish

results of testing the agar with iodine solution after incubation with halves of maize grains

Fig. 1.1

The students thought that the area stained brown was proportional to the activity of the amylase and could be used to test the hypothesis:

The greater the concentration of gibberellin (GA) to which the maize is exposed, the greater the activity of amylase.

dentify the independent and the dependent variables in the students' preliminary nvestigation.				
dependent variable				
ependent variable				
The students were provided with a 3 mmol dm ⁻³ Describe how the students could use the method test their hypothesis. Your method should be couse.	from their preliminary investigation detailed enough for another person			

.....[8]

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(b)	The students decided to plot a graph of their results.					
Suggest labels, including units, for the axes of this graph.						
	x-axis					
	<i>y</i> -ax	dis				
		[2]				
(c)	(i)	The students thought that the area stained brown was proportional to the activity of the amylase.				
		Suggest three limitations of using this way to estimate amylase activity.				
		1				
		2				
		3				
		[3]				
	(ii)	For one of these limitations, suggest how the estimation of amylase activity could be improved.				
		[2]				
		[Total: 17]				

2 In an area where an open cast or surface mine was to be developed, the soil was removed and stored in a large heap. After the mining was finished, the area was reclaimed and the soil spread over the surface.

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Studies were carried out on the aerobic and anaerobic bacteria living in the soil while it was stored.

Samples were taken at different depths, 1 month and 6 months after the soil was put into the large heap for storage.

Table 2.1 shows the numbers of aerobic and anaerobic bacteria at different depths in the stored soil.

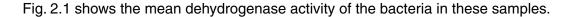
Table 2.1

	mean number of bacteria per gram of stored soil \times 10 ⁷				
depth in soil store / m	aerobic	bacteria	anaerobic bacteria		
	after 1 month	after 6 months	after 1 month	after 6 months	
0.0	12.4	12.5	0.4	0.6	
0.5	10.1	8.3	0.6	1.0	
1.0	9.8	5.9	0.8	3.8	
1.5	9.7	3.1	0.8	7.6	
2.0	10.5	0.8	0.7	8.1	
2.5	10.8	0.7	0.8	8.5	
3.0	10.2	0.9	0.6	8.8	

(a)	Describe the trends shown by the distribution of the two types of bacteria in the stored soil after 6 months and suggest reasons for these trends.
	[3]

(b) In a further study, soil samples were taken at two depths, **A** and **B**, in the soil store. The samples were taken at intervals over six years. Soil samples of equal mass were used to determine the activity of dehydrogenase enzymes in the Krebs cycle of the **aerobic** bacteria.

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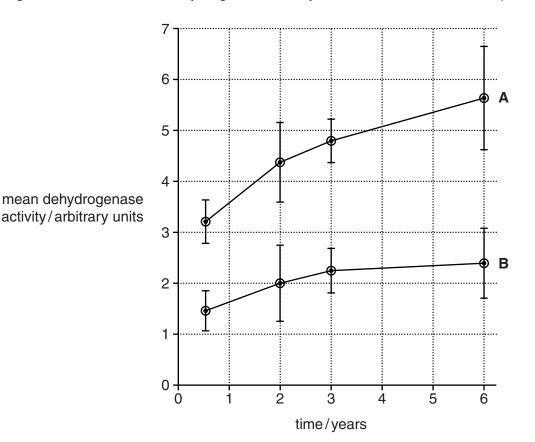


Fig. 2.1

(i)	State the evidence from Fig. 2.1 that samples in ${\bf B}$ were taken at a greater depth than samples in ${\bf A}$.
	[1]
(ii)	Suggest two variables that should be standardised when the dehydrogenase activity is determined.
	1
	2
(iii)	Suggest a control for the determination of dehydrogenase activity in the bacteria in the soil samples.
	[1]

(c)	The	The error bars on Fig. 2.1 are two standard error units above and below the mean.						
	(i)	(i) Using evidence from Fig. 2.1 state what these error bars show about the roof the data.						
(ii) A t-test was carried out between each of the following sets of data from th at depth A:				of data from the samples				
		6 months and 2 years;	6 months and 3 years;	6 months and 6 years;				
		2 years and 3 years;	2 years and 6 years;	3 years and 6 years.				
		Identify two <i>t</i> -tests in which significant. Show your answ	n the difference in dehydrogena ver by underlining the tests.	ase activity is likely to be [1]				
		Give a reason for your answ	ver.					
				[1]				
(d)		le 2.2 shows the dehydroger ix soil samples.	nase activity and the number of	aerobic bacteria present				
			Table 2.2					
dehy	/drog	enase activity / arbitrary unit	s number of aerobic bacter	ia per gram of soil \times 10 ⁷				
		13.5	12.	2				
		9.6	9.	1				
		5.8	7.	0				
		3.0	4.	6				
		2.5	3.	2				
		0.6	0.	9				
	Exp		g of another soil sample was no could be used to predict the nu					
				[2]				

[Total: 13]

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