

Cambridge Pre-U

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		

BIOLOGY 9790/02

Paper 2 Data Analysis and Planning

For examination from 2020

SPECIMEN PAPER

1 hour 15 minutes

You must answer on the question paper.

No additional materials are needed.

INSTRUCTIONS

- Answer all questions.
- Use a black or dark blue pen. You may use an HB pencil for any diagrams or graphs.
- Write your name, centre number and candidate number in the boxes at the top of the page.
- Write your answer to each question in the space provided.
- Do not use an erasable pen or correction fluid.
- Do not write on any bar codes.
- You may use a calculator.
- You should show all your working and use appropriate units.

INFORMATION

- The total mark for this paper is 60.
- The number of marks for each question or part question is shown in brackets [].

This specimen paper has been updated for assessments from 2020. The specimen questions and mark schemes remain the same. The layout and wording of the front covers have been updated to reflect the new Cambridge International branding and to make instructions clearer for candidates.

For Examiner's Use		
Section A		
Section B		
Total		

This syllabus is regulated for use in England, Wales and Northern Ireland as a Cambridge International Level 3 Pre-U Certificate.

This document has 18 pages. Blank pages are indicated.

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Answer all the questions.

Section A - Data analysis

1 Fig. 1.1 shows an American eel, *Anguilla rostrata*, which lives for part of its life in the rivers and mountain streams of the Eastern USA. Adult fish migrate to the Atlantic Ocean when they are ready to breed. After breeding the adults die.

Young eels migrate from the sea back to the rivers and streams and may live for five years or more before reaching the stage when they are ready to breed.

This species of fish has become rare in mountain streams over recent years.



Fig. 1.1

As part of a long-running study to find out more about the biology and behaviour of *A. rostrata*, mark-release-recapture was used to estimate the population size in one mountain stream in Virginia. Very young eels were not marked.

Table 1.1 shows the results of the mark-release-recapture.

Table 1.1

year	total number of fish caught	number of fish marked	total number of fish captured one year later	number of marked fish recaptured one year later	population estimate
2000	334	279	352	98	1002
2001	352	226	290	57	1150
2002	290	149	180	25	1073
2003	180	76	232	11	1603
2004	232	116	184	21	

(a) (i) Complete Table 1.1 by calculating the population estimate for the year 2004.

(ii)	Comment on the accuracy of the population estimates for eels recorded in Table 1.1, including limitations of the mark-release-recapture method for estimating population size.

[1]

The annual growth of the eels was also measured. Fig. 1.2 shows a box-whisker plot of the results for growth in length and growth in mass of eels in one stream that were marked with tags and then recaptured from 2000 to 2005.

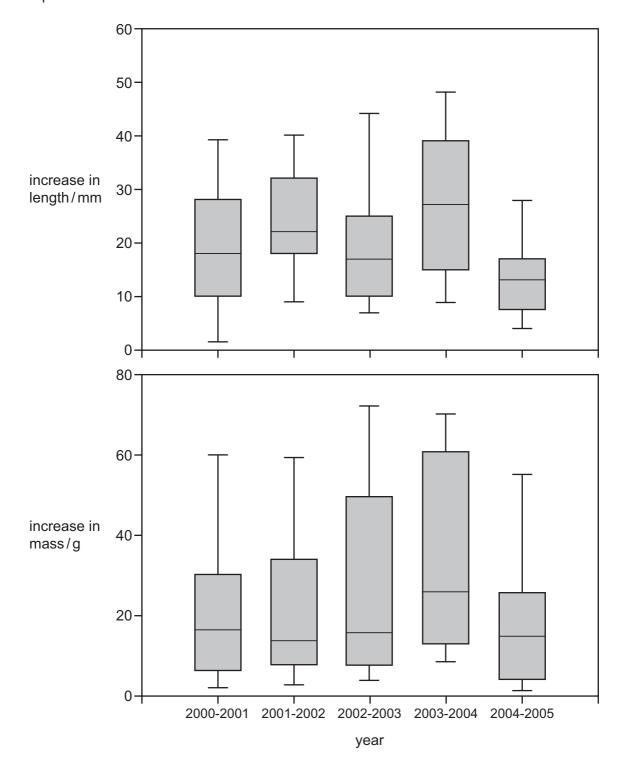


Fig. 1.2

The horizontal line in each box represents the median. The top and bottom of each box show 25th and 75th percentiles. The 'whiskers' show the 10th and 90th percentiles.

(b)	(i)	Describe the results shown in Fig. 1.2.
		[3]
	(ii)	Suggest advantages of using box-whisker plots to show these data rather than bar charts or histograms.
		[3]

 A. rostrata is not officially recognised as an endangered species, but its numbers are in decline.
Discuss the limitations of the results of this study in terms of providing sufficient information to inform the conservation of <i>A. rostrata</i> in Virginia.
[4
[Total: 17]

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In an investigation into pollen release from Timothy grass, *Phleum pratense*, the number of pollen grains released into the atmosphere was sampled at hourly intervals, on three consecutive days. Traps sited just above the level of the leaves were used to do this.

The wind speed and the relative humidity were recorded at the times of sampling.

The results of the investigation are shown in Fig. 2.1.

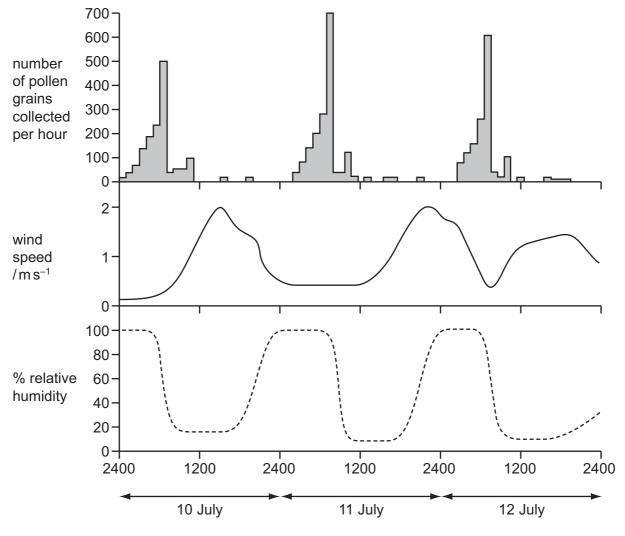


Fig. 2.1

(a)	Describe the results shown in Fig. 2.1.
	[6]
(b)	One researcher concluded that high humidity and low wind speed were important factors for the release of pollen from Timothy grass.
	Discuss why this conclusion may not be valid.
	[4]
	[1]

[Total: 10]

The technique of polyacrylamide gel electrophoresis (PAGE) is used to separate and identify proteins. One method of PAGE involves treating proteins with an ionic detergent to dissociate proteins into their constituent polypeptide subunits. Sodium dodecyl sulfate (SDS) is often used for this. Proteins treated with SDS have a uniform net charge on each polypeptide so that during electrophoresis they are separated only on the basis of their relative molecular mass.

After treatment with SDS, proteins are placed in wells cut into a polyacrylamide gel. A dye is added to each sample to show the progress of the samples across the gel. A current is applied to the gel and when the dye reaches a point towards the end of the gel, the current is switched off.

The relative mobility of each polypeptide is calculated as follows:

distance travelled by polypeptide band distance travelled by dye front

Six proteins, **A**, **B**, **C**, **D**, **E** and **F**, were analysed with SDS-PAGE and the results are shown in Fig. 3.1.

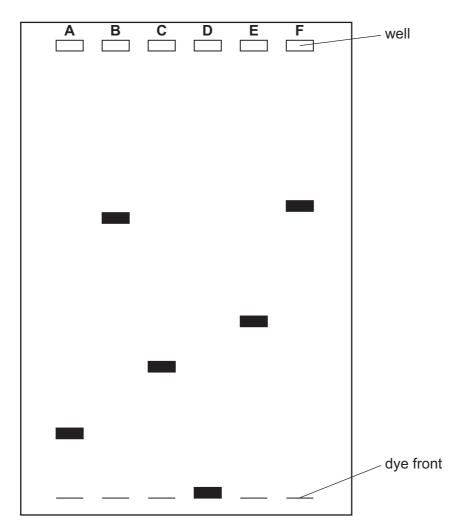


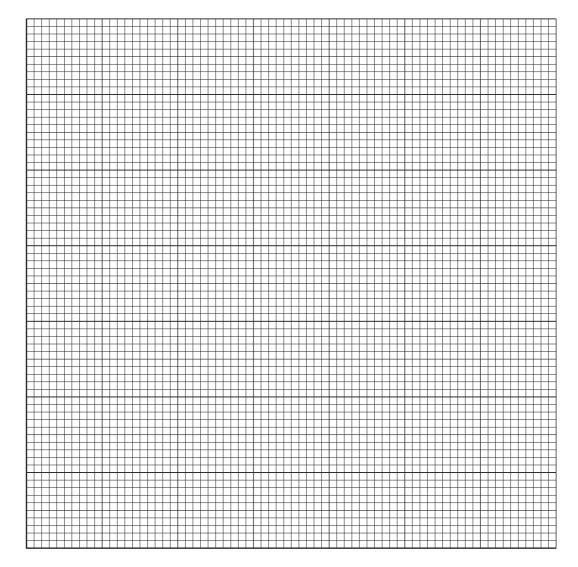
Fig. 3.1

(a) (i) Calculate the relative mobility of proteins A, B, C, E and F and add your calculated values to the appropriate spaces in Table 3.1, opposite. [2]

Table 3.1

protein	relative molecular mass	relative mobility
Α	29 000	
В	68 000	
С	unknown	
D	17200	1.00
E	43 000	
F	77 000	

(ii) Plot, on the grid below, a graph of the relative molecular mass of proteins **A**, **B**, **D**, **E** and **F** against their relative mobility. [4]



(b)	Use your graph to find the relative molecular mass of protein C .
	Explain how you arrived at your answer.
	relative molecular mass of protein C:
	[2
	•

[Total: 8]

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Section B - Planning

4 Yeast cells have transport proteins in their cell membranes for the uptake of nutrients from the surroundings. There are separate transport proteins for glucose and for maltose. When exposed to both glucose and maltose the transport protein for maltose is downregulated and is not produced.

Plan an investigation to find out whether or not the yeast transport proteins for glucose and maltose function at the same rate.

Glucose and maltose are both reducing sugars.

You are provided with the following materials. Choose your materials from this list. You may **not** use any additional materials.

- 10% yeast suspension
- 10 g dm⁻³ glucose solution 10 g dm⁻³ maltose solution
- Benedict's solution
- dilute hydrochloric acid
- dilute sodium hydroxide solution and sodium hydrogencarbonate solution for neutralising
- beakers and flasks of different sizes
- stopwatch or electronic timer
- colorimeter and tubes
- centrifuge and centrifuge tubes
- thermometer
- thermostatically-controlled water baths
- pipettes and pipette fillers
- burettes and burette stands
- filter funnels and filter paper
- svringes
- glass rods for stirring
- test-tubes and boiling tubes
- test-tube and boiling tube racks

Your plan should

- include a clear statement of the hypothesis or prediction
- identify the key variables
- give full details and explanations of the procedures that you would adopt to ensure that the results are as precise and repeatable as possible
- show how you would present and analyse your results
- include a brief risk assessment
- be written in clear, scientific language.

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Copyright Acknowledgements:

Question 1 Figure 1.1 American Eel © Andrew J. Martinez/Science Photo Library

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