| Write your name here Surname | Oth | er names |
|--|---------------|--------------------------|
| Edexcel GCE | Centre Number | Candidate Number |
| Biology Advanced Subsidi Unit 4: The Natura Survival | • | and Species |
| Wednesday 16 June 2010 Time: 1 hour 30 minute | • | Paper Reference 6BI04/01 |
| You do not need any other | materials. | Total Marks |

Instructions

- Use **black** ink or ball-point pen.
- **Fill in the boxes** at the top of this page with your name, centre number and candidate number.
- Answer **all** questions.
- Answer the questions in the spaces provided
 - there may be more space than you need.

Information

- The total mark for this paper is 90.
- The marks for **each** question are shown in brackets
 - use this as a guide as to how much time to spend on each question.
- Questions labelled with an asterisk (*) are ones where the quality of your written communication will be assessed
 - you should take particular care with your spelling, punctuation and grammar, as well as the clarity of expression, on these questions.
- Candidates may use a calculator.

Advice

- Read each question carefully before you start to answer it.
- Keep an eye on the time.
- Try to answer every question.
- Check your answers if you have time at the end.

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Answer ALL questions.

Some questions must be answered with a cross in a box \boxtimes . If you change your mind about an answer, put a line through the box \boxtimes and then mark your new answer with a cross \boxtimes .

- **1** Forensic scientists use several different types of evidence to estimate the time of death of a body.
 - (a) Place a cross ⋈ in the boxes next to the **two** types of evidence that could be used by a forensic scientist to estimate the time of death of a body.

(2)

- Degree of muscle contraction
- □ Length of fingernails
- Signs of decomposition
- Skin pigmentation
- (b) The core temperature of a body can also be used to estimate the time of death. This can be recorded by measuring the temperature of the rectum.

The table below shows the mean core temperature, measured at 2-hour intervals after death, using data from a large number of bodies. The standard deviation from each of the means is also shown.

| Time after death / hours | Mean core temperature /°C | Standard deviation from the mean / °C |
|-----------------------------|------------------------------|---------------------------------------|
| 2 | 36.7 | 0.9 |
| 4 | 36.0 | 1.1 |
| 6 | 35.3 | 1.3 |
| 8 | 31.3 | 1.5 |
| 10 | 29.6 | 1.7 |
| 12 | 28.1 | 2.0 |
| 14 | 26.7 | 2.3 |
| 16 | 25.5 | 2.6 |
| 18 | 25.0 | 2.9 |
| 20 | 23.6 | 3.1 |
| 22 | 22.8 | 3.3 |
| 24 | 22.3 | 3.4 |
| 26 | 21.8 | 3.4 |

| Suggest three factors that could influence the rate at which a body cools after death. (Total for Question 1 = 9 marks) | | |
|--|--|--------------|
| death. (3) | | |
| | Suggest three factors that could influence the rate at which a body cool death. | s after |
| (Total for Question 1 = 9 marks) | | (3) |
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| 2 | It has been estimated that only 5% of the light energy hitting the surface of a leaf | |
|---|---|-----|
| _ | reaches the chloroplasts to be used in the synthesis of organic material. The total energy used in this synthesis is known as the gross primary productivity (GPP). | |
| | (a) Suggest two reasons why 95% of the light hitting the surface of a leaf is not used by the chloroplasts. | |
| | | (2) |
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| | | |
| | (b) The mean GPP for plants on Earth is 24.4×10^6 J m ⁻² year ⁻¹ . | |
| | The plants use 3.7×10^6 J m ⁻² year ⁻¹ of this energy in metabolic processes. The energy in the remaining organic material is known as net primary productivity (NPP). | |
| | (i) Explain what is meant by the unit J m⁻² year⁻¹ . | (1) |
| | (ii) Calculate the percentage of the mean GPP that remains as NPP within plants on Earth. | |
| | Show your working. | (2) |
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| | | |
| | Angwar | % |
| | Answer | 90 |
| | | |
| | | |

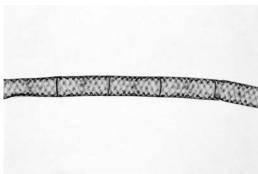
| X | | | (1) |
|---|---|--|---------|
| | A | Chemosynthesis | |
| X | В | Respiration | |
| X | C | Photosynthesis | |
| X | D | Protein synthesis | |
| | | ence to the structures in a chloroplast, explain how the energy from de available in ATP molecules for the synthesis of organic materials. | (6) |
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| | | (Total for Question 2 = 12 ma | - ulaa) |

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3 Filamentous algae are simple photosynthetic organisms that consist of long strands of very similar eukaryotic cells. Each of the cells in the strand is enclosed within a cellulose cell wall. The strand increases in length as the cells divide and elongate.

The photographs below show some cells in strands of a filamentous alga, as seen using a light microscope.





Magnification ×200

Magnification ×200

(a) (i) Put a cross ⊠ in the box next to the term that describes the process involved in the cell divisions in a filamentous alga.

(1)

- A exocytosis
- **B** meiosis
- **C** mitosis
- **D** osmosis
- (ii) Put a cross ⊠ in the box next to the structure that would **not** be found in a cell from the strand of a filamentous alga.

(1)

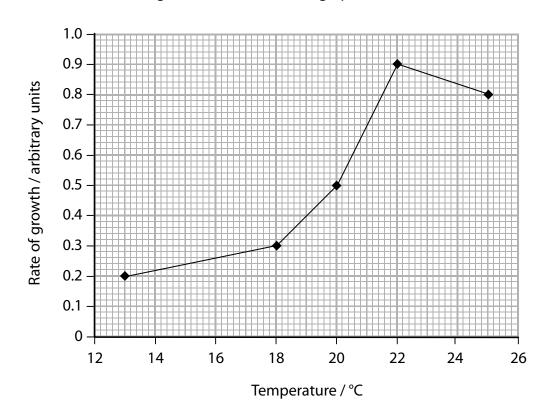
- A lysosome
- **B** mitochondrion
- **C** plasmid
- **D** ribosome
- (b) An investigation was carried out into the effect of temperature on the rate of growth of a filamentous alga. Several short strands of the alga were placed into culture solutions which were kept at five different temperatures and at a high light intensity.

The number of cells in the strands, in each culture solution, was counted at the beginning of the time period and again after 18 days.

The rate of growth was then calculated.



The results of this investigation are shown in the graph below.



(i) Name the **independent** variable in this investigation.

(1)

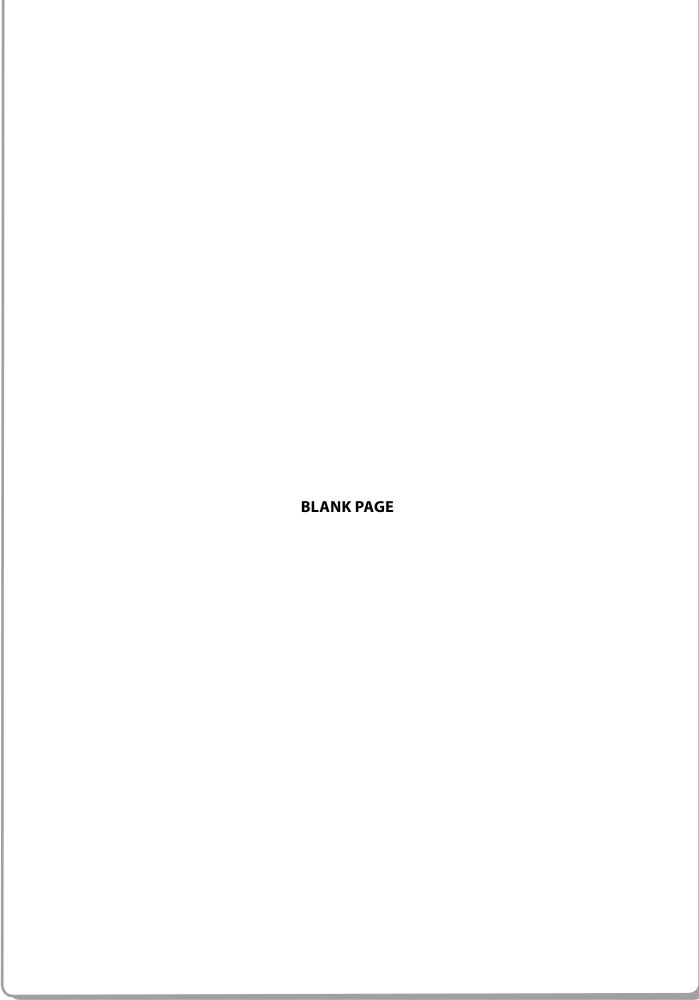
(ii) Using the information in the graph, describe and suggest explanations for the effect of temperature on the rate of growth of the filamentous alga.

(4)



| | light intensity. (3) |
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| iv) | Suggest two abiotic factors, other than light intensity, that would need to be controlled in this investigation. |
| | (2) |
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| | (Total for Question 3 = 12 marks) |
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| 4 Various non-specific responses of the body are involved in resistance to bacterial infections. (a) Histamine, interferon and lysozyme are three substances involved in non-specific responses to infection. For each of the following statements, put a cross ⋈ in the box next to the name of the substance involved. (i) Enzyme released in secretions that break down the cell walls of bacteria. A Histamine B Interferon C Lysozyme (ii) Inflammation caused by a chemical released by white cells in connective tiss A Histamine B Interferon C Lysozyme | |
|--|------------|
| responses to infection. For each of the following statements, put a cross in the box next to the name of the substance involved. (i) Enzyme released in secretions that break down the cell walls of bacteria. A Histamine B Interferon C Lysozyme (ii) Inflammation caused by a chemical released by white cells in connective tiss A Histamine B Interferon | - 1 |
| the substance involved. (i) Enzyme released in secretions that break down the cell walls of bacteria. A Histamine B Interferon C Lysozyme (ii) Inflammation caused by a chemical released by white cells in connective tiss A Histamine B Interferon | |
| ■ A Histamine ■ B Interferon ■ C Lysozyme (ii) Inflammation caused by a chemical released by white cells in connective tiss ■ A Histamine ■ B Interferon | f |
| ■ B Interferon ■ C Lysozyme (ii) Inflammation caused by a chemical released by white cells in connective tiss ■ A Histamine ■ B Interferon | (1) |
| ■ C Lysozyme (ii) Inflammation caused by a chemical released by white cells in connective tiss ■ A Histamine ■ B Interferon | · I— |
| (ii) Inflammation caused by a chemical released by white cells in connective tiss ■ A Histamine ■ B Interferon | |
| ■ A Histamine■ B Interferon | |
| ■ B Interferon | ue. (1) |
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| ☑ C Lysozyme | |
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| | Bacteria with D on surface enter blood |
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| | E produce F which bind to D and label bacteria |
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| | Labelled bacteria attach to F receptors on G |
| | |
| | G engulfs labelled bacteria by forming vacuoles |
| | |
| | H is released into the vacuoles to destroy the bacteria |
| (i) Identify D , E | E , F , G and H by writing appropriate terms in the spaces below. (5) |
| D | |
| E | |
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| response to some types of bacte | ria. |
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| | (3) |
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| Description | DNA only | RNA only | Both DNA and RNA | |
|--|--|---|---|--|
| Polymer formed from a single strand of nucleotides | | | | |
| Pentose present in the nucleotides | | | | |
| Adenine, cytosine, guanine and thymine present | | | | |
| Nucleotides linked by phosphodiester bonds | | | | |
| - | | lowing proce | sses leads to t | he |
| esis of this sequence of amino ac | cids. | | | |
| ne formation of mkNA during tr | anscription ir | i the nucleus | | (3) |
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| | | | | |
| ֡ | Polymer formed from a single strand of nucleotides Pentose present in the nucleotides Adenine, cytosine, guanine and thymine present Nucleotides linked by phosphodiester bonds liagram below shows the sequencule. RNA anticodon that corresponds o acids Alanine—Glutam anticodon CGA GUU gethis information, explain how explain sequence of amino acids of this sequence of amino acids and control of the sequence of amino acids of this sequence of t | Polymer formed from a single strand of nucleotides Pentose present in the nucleotides Adenine, cytosine, guanine and thymine present Nucleotides linked by phosphodiester bonds liagram below shows the sequence of the last cule. RNA anticodon that corresponds to each amino acids a cids Alanine—Glutamine—Glycin anticodon CGA GUU CCA | Polymer formed from a single strand of nucleotides Pentose present in the nucleotides Adenine, cytosine, guanine and thymine present Nucleotides linked by phosphodiester bonds liagram below shows the sequence of the last six amino accule. RNA anticodon that corresponds to each amino acid is also acids a acids Alanine—Glutamine—Glycine—Aspara anticodon CGA GUU CCA UU/ g this information, explain how each of the following processes of this sequence of amino acids. | Polymer formed from a single strand of nucleotides Pentose present in the nucleotides Adenine, cytosine, guanine and thymine present Nucleotides linked by phosphodiester bonds liagram below shows the sequence of the last six amino acids in a proteicule. RNA anticodon that corresponds to each amino acid is also shown. o acids Alanine—Glutamine—Glycine—Asparagine—Prolin anticodon CGA GUU CCA UUA GGA |

| (ii) The translation of mRNA into the sequence of amino acids in a ribosome | (3) |
|---|------|
| | |
| (c) Suggest why the final triplet of nucleotides, on the strand of mRNA involved in the synthesis of this sequence of amino acids, did not correspond with any anticodon on tRNA. | |
| | (2) |
| | |
| (Total for Question 5 = 10 mar | ·ks) |
| | |
| | |
| synthesis of this sequence of amino acids, did not correspond with any anticodon on tRNA. | (2) |

| б | Rhododendrons are shrubby plants that are widely distributed throughout the |
|---|---|
| | northern hemisphere. |

The flowering periods and habitats of two species of rhododendron, found on Yakushima Island in Japan, are shown in the table below.

| Species | Flowering period | Main flowering period | Habitat |
|-------------------------|------------------|-----------------------|--------------------------------|
| Rhododendron eriocarpum | April to July | May | Rocky areas in lowland regions |
| Rhododendron indicum | May to July | June | High mountainous regions |

Where these populations overlap, hybrid plants are found that have arisen as a result of cross-fertilisation between these two species. These hybrid plants are capable of flowering and producing viable seeds.

| separate species. | | | (3) |
|------------------------------------|--------------------------------|-------------------------|-----|
| | | | (3) |
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|) (1) F 1: 1 : | | | |
|) (i) Explain what is meant by the | e term genetic divers i | ty in a species. | (2) |
| | | | (2) |
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| (ii) Explain why there is likely to be a greater genetic diversity in the hybrid plant than in either of the two separate species. | s |
|--|------|
| | (2) |
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| | |
| *(c) Explain how the two different species of Rhododendron on Yakushima Island may have evolved from a single population of an ancestral species. | |
| have evolved from a single population of an ancestral species. | (6) |
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| (Total for Question 6 = 13 ma | rks) |
| (Total for Question 0 – 13 ma | 113/ |



| 7 | Several species of woodlice can be found in most gardens in Britain. Woodlice have |
|---|---|
| | bacteria in their digestive system that secrete enzymes required for the digestion of |
| | plant cell walls. Woodlice are decomposers of dead plants. |

The photograph below shows one of the common species of woodlouse found in gardens in Britain.



Magnification x10
Amateur Entomologists Society / Kieren Pitts

| (a) Suggest how woodlice are involved in the recycling of carbon. | (3) |
|---|-----|
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| at f | a of paving slabs were fairly active and running about. When she lifted some of stones near the slabs, the woodlice under these stones were relatively inactive first. However, within a very short time these woodlice started to run about. For a few minutes, almost all of the woodlice that were under the stones had | |
|------|---|-----|
| | appeared from view. | |
| (i) | The student thought that the behaviour and distribution of the woodlice were being influenced by an abiotic factor in her garden. | |
| | Place a cross \boxtimes in the box next to the term that describes this type of idea. | (1) |
| X | A Hypothesis | |
| X | B Observation | |
| X | C Prediction | |
| X | D Theory | |
| (ii) | Suggest two examples of abiotic factors that might influence the behaviour and distribution of the woodlice in her garden. | |
| | | (2) |
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(c) She decided to investigate the behaviour of the woodlice. She collected ten woodlice and released them into the centre of the area of paving slabs. She took photographs immediately after the woodlice were released (time 0) and each minute for ten minutes. This was repeated twice. Using the photographs, she recorded the number of woodlice on the paving slabs at one-minute intervals.

Her results are shown in the table below.

| Time after | Number of woodlice on paved area | | | |
|----------------------|----------------------------------|-------------|-------------|------|
| release / minutes | 1st release | 2nd release | 3rd release | Mean |
| 0 | 10 | 10 | 10 | 10 |
| 1 | 10 | 9 | 9 | 9 |
| 2 | 9 | 8 | 9 | 9 |
| 3 | 7 | 7 | 7 | 7 |
| 4 | 7 | 6 | 5 | 6 |
| 5 | 6 | 6 | 6 | 6 |
| 6 | 4 | 5 | 4 | 4 |
| 7 | 4 | 3 | 4 | 4 |
| 8 | 3 | 3 | 2 | |
| 9 | 0 | 2 | 1 | |
| 10 | 1 | 1 | 0 | |

(i) Complete the table by calculating the mean for each of the final three minutes.

(1)



| (ii) | Suggest why taking photographs is a suitable method to count the woodlice. | (2) |
|-------|--|-----|
| | | |
| (iii) | Explain why it would be difficult to determine which abiotic factor is influencing the behaviour and distribution of the woodlice in a garden environment. | (3) |
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| | (Total for Question 7 = 12 mar | ks) |
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| | | (3) |
|--|-----------------------|-----------------|
| Description | Name of structure | P, E or B |
| Enclosed by outer smooth membrane; inner membrane folded forming cristae | | |
| Long strand-like structure extending out from the cell; used for locomotion | | |
| Small, circular loop of double-stranded DNA | | |
| peptidoglycan, a component of l | | |
| peptidoglycan, a component of logical (i) State the term used to descress bacterial cells. | oacterial cell walls. | ycin, that kill |
| | oacterial cell walls. | ycin, that kill |
| peptidoglycan, a component of l (i) State the term used to descr bacterial cells. | oacterial cell walls. | ycin, that kill |
| peptidoglycan, a component of l (i) State the term used to descr bacterial cells. | oacterial cell walls. | ycin, that kill |
| peptidoglycan, a component of l (i) State the term used to descr bacterial cells. | oacterial cell walls. | ycin, that kill |
| peptidoglycan, a component of logical (i) State the term used to descress bacterial cells. | oacterial cell walls. | ycin, that kill |

Antibiotics are used to treat bacterial infections in eukaryotic organisms.



| (iii) Explain why doctors have been advised to limit the prescription of antibiotics. (2) | |
|--|--|
| | |
| (c) Describe how you could investigate the effect of different antibiotics on bacteria. (4) | |
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| (Total for Question 8 = 12 marks) | |
| TOTAL FOR PAPER = 90 MARKS | |
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