Please write clearly in	plock capitals.
Centre number	Candidate number
Surname	
Forename(s)	
Candidate signature	

A-level BIOLOGY

Unit 5 Control in cells and in organisms

Thursday 23 June 2016

Morning

Time allowed: 2 hours 15 minutes

Materials

For this paper you must have:

- a ruler with millimetre measurements
- a calculator.

Instructions

- Use black ink or black ball-point pen.
- Fill in the boxes at the top of this page.
- Answer all questions.
- You must answer the questions in the spaces provided. Do not write outside the box around each page or on blank pages.
- You may ask for extra paper. Extra paper must be secured to this booklet.
- Do all rough work in this book. Cross through any work you do not want to be marked.

Information

- The marks for questions are shown in brackets.
- The maximum mark for this paper is 100.
- You are expected to use a calculator, where appropriate.
- Quality of Written Communication will be assessed in all answers.
- You will be marked on your ability to:
 - use good English
 - organise information clearly
 - use scientific terminology accurately.

Advice

You are advised to spend no longer than 40 minutes on the essay.



	Answer all questions in the spaces provided.
1 (a)	 The following statements are about events during an action potential. A Potassium ions diffuse out across the neurone membrane. B Sodium ions diffuse in across the neurone membrane. C Sodium ion channels open. D Active transport of sodium and potassium ions restores resting potential. E Potassium ion channels open. F Hyperpolarisation of the membrane occurs.
1 (a) (i)	Which of the events, A to F , starts depolarisation? Put the correct letter in the box. [1 mark]
1 (a) (ii)	Which of the events, A to F , requires the hydrolysis of ATP? Put the correct letter in the box. [1 mark]
1 (b)	Synaptophysin is a protein involved in the production of synaptic vesicles.
	Scientists can use the presence or absence of synaptophysin to identify presynaptic and postsynaptic membranes in synapses.
	Explain why they are able to use synaptophysin for this purpose. [1 mark]



1 (c) Dopamine is a neurotransmitter. Production of too much dopamine is associated with schizophrenia. A drug used to treat schizophrenia binds to dopamine receptors in synapses. This binding does not lead to the formation of an action potential. 1 (c) (i) Suggest why the drug used to treat schizophrenia is able to bind to the same receptor as dopamine. [1 mark] 1 (c) (ii) Suggest why binding of the drug does not lead to production of an action potential. [2 marks] Turn over for the next question



2 (a)	It is important that mammals maintain a constant core temperature.
	Explain why. [3 marks]
2 (b)	Scientists investigated control of body temperature in rats. Rats have two patches of brown adipose tissue (BAT) under the skin of their shoulders. In BAT, respiration takes place in mitochondria but no ATP is formed. The scientists looked at the response of BAT to cooling of the rat's skin.
	 The scientists cooled and warmed an area of a rat's skin. At the same time, they recorded the following. Electrical activity in neurones that control the activity of BAT. The temperature of the BAT.
	 The percentage of carbon dioxide in the rat's breath. Figure 1 shows their results.





Turn over ►



Impala and wildebeest are species of herbivore that live in large groups. They spend most of their time feeding with their heads near the ground.

Scientists investigated the relationship between the number of predators in an area and the mean proportion of time these herbivores spent with their heads up, looking around rather than feeding. They obtained data from groups of impala and wildebeest in two areas. In one area there were few predators and in the other area there were many predators.





[2 marks]		Show your working.
nce hours	Difference	



3 (b)	The scientists concluded that these herbivores spend more time looking for predators in areas where there are many predators.
	Do these data support this conclusion? Give reasons for your answer. [4 marks]
3 (c)	The behaviour of the herbivores in having their heads up has a benefit but it also has costs. The benefit is being able to see, and escape from, predators.
	Suggest and explain one cost to the herbivores of this behaviour. [2 marks]
	Turn over for the next question







4 (b) (i)	What was the percentage fall in the mean force produced by mice not able to produce creatine, compared with the normal mice? Show your working. [2 marks]
	Answer %
4 (b) (ii)	Suggest an explanation for these results. [2 marks]
4 (c)	The mice that were not able to produce creatine were homozygous for a recessive allele
	of a gene. Mice that are heterozygous for this allele are able to produce forces similar to those of normal mice that are homozygous for the dominant allele of the same gene. Explain why the heterozygous mice can produce forces similar to those of normal mice. [2 marks]
	Turn over for the next question



Agrobacterium tumefaciens is a bacterium that is often used in recombinant DNA technology to produce transformed plants that benefit humans.

A. tumefaciens contains a plasmid which can be used as a vector to transfer a desired gene into plant cells. These plant cells may then develop into plants which produce the protein coded for by the desired gene.

Figure 4 outlines this process.





[1 mark]

In stage 1, an enzyme is used to cut open the plasmid. Name the type of enzyme used to cut open the plasmid.

5 (a) (i)

5 (a) (ii)	In stage 1, another enzyme is used to insert the desired gene into the plasmid DNA. Name the type of enzyme used to insert the gene into the plasmid. [1 mark]
5 (b)	In stage 4, some plant cells had plasmid DNA only in their cytoplasm. In other plant cells, the plasmid DNA had become inserted into plant DNA in the nucleus.
	In stage 5, only cells with plasmid DNA inserted into the plant DNA in the nucleus grew into plants where all the cells contained the desired gene.
	Explain why some of the plants in stage 5 contained the desired gene in all of their cells and others did not.
	[3 marks]
5 (c)	The desired gene in Figure 4 was from an insect. In stage 6, the plant containing this gene was able to use it to synthesise an insect protein.
	The plant is able to synthesise the insect protein. Explain why this is possible. [3 marks]
	Turn over ▶
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over 🕨













7 (a)	Use information in Figure 6 to suggest how the binding of interferon gamma receptor protein leads to the production of phosphorylated STAT1.	to its
	ere han ere ere ere ere ere ere ere ere ere er	[2 marks]
7 (b)	Name the two transcription factors in Figure 6 .	[2 marks]
	1	
	2	
7 (c)	The regulation of the formation of helper T cells by interferon gamma is an expositive feedback.	
	Explain why it is an example of positive feedback.	
		[2 marks]
7 (d)	The <i>IRF</i> gene can be a tumour suppressor gene.	
	Use the information in Figure 6 to explain how the <i>IRF</i> gene acts as a tumou	r
	suppressor gene.	[3 marks]



8	<i>Mycobacterium tuberculosis</i> causes tuberculos a direct repeat (DR) region. The DR region con sequences called spacers. Each spacer is four In different strains of <i>M. tuberculosis</i> , some of	sists of 43 different, non-coding base nd in a specific place in the DR region.
8 (a) (i)	The DR region consists of non-coding base see	quences.
	What is meant by a non-coding base sequence	? [1 mark]
8 (a) (ii)	Name the process by which the base sequence	e of a spacer is lost from a DR region. [1 mark]
	 Scientists investigated the DR regions of difference produced a DNA probe for each of the 43 space Iabelled with a fluorescent marker that gave complementary spacer attached to a particular square on a slide. They obtained samples of the DR region from existingle-stranded DNA fragments. The fragment with the DNA probes attached. Figure 7 shows <i>M. tuberculosis</i> with 20 of the probes. 	er sequences. Each probe was: off light if the probe attached to its each strain. These were cut into small s from each strain were added to a slide
	rigure /	
Slide DR fr addeo		Square where no light was seen (black square)



8 (b)	The scientists cloned the DR region DNA <i>in vitro</i> before testing for the presence of spacers.
	Give the name of the method they used to clone the DNA <i>in vitro</i> . [1 mark]
8 (c)	Explain how the use of DNA probes produced the results in Figure 7 . [3 marks]
8 (d)	Doctors can use the method with DNA probes to identify the specific strain of <i>M. tuberculosis</i> infecting a patient. This is very important when there is an outbreak of a number of cases of tuberculosis in a city.
	Suggest and explain why it is important to be able to identify the specific strain of <i>M. tuberculosis</i> infecting a patient.
	[2 marks]
	Turn over for the next question



9 Multiple sclerosis (MS) is a condition caused when the body's own immune system attacks the myelin sheath around axons. The cell bodies of the neurones themselves can also be damaged or destroyed. People with MS usually have periods of time when their MS gets no worse, followed by relapses when it gets worse.

Scientists investigated the effects on neurones of damage to myelin. The scientists obtained a modified antigen from the myelin sheath of humans and injected it into mice. After a number of days, this injection of antigen resulted in the myelin sheaths in the mice being damaged. Some cell bodies of neurones were also damaged.

9 (a) Suggest how the injection of the antigen resulted in the myelin sheaths being damaged. [3 marks]

9 (b) The scientists compared the ultrastructure of normal and damaged neurones. They found that damaged neurones contained many mitochondria with an unusual ultrastructure.

Figure 8 shows a mitochondrion with normal ultrastructure and one with the unusual ultrastructure.

Figure 8





	Suggest why having a large number of mitochondria with this unusual ultrastructure could lead to neurones dying. [3 marks]
9 (c)	The scientists took a large number of photographs of thin sections through neurones. Using these photographs, they found that 40% of mitochondria had the unusual ultrastructure in damaged neurones.
9 (c) (i)	What sort of microscope would the scientists use to take the photographs? Give one reason for your answer. [1 mark]
	Type of microscope
	Reason
9 (c) (ii)	Suggest how the scientists found the percentage of mitochondria with the unusual ultrastructure.
	[3 marks]



9 (d)	treat MS. They recruited into three groups, A , B a	sts investigated the use of a drug called teriflunomide to a large number of volunteers who had MS and divided them and C , at random. For each group, they recorded factors such ses they had and how long it was since they were diagnosed
9 (d) (i)	Explain why the scientist	s made these comparisons. [1 ma
9 (d) (ii	i) Each group of volunteers given to each group was	s was given a different treatment for 2 years. The treatment as follows.
	Group B was given 7	placebo that contained no drug. mg of teriflunomide per day. 4 mg of teriflunomide per day.
	The scientists determine group.	d the mean number of relapses per person, per year for each
	Table 1 shows their resu	ilts.
		Table 1
	Group	Mean number of relapses of MS per person per year (± 95% confidence limits)
	A (placebo)	0.55 (± 0.10)
	B (7 mg teriflunomide)	0.37 (± 0.07)
	1	0.36



Evaluate this conclusion.	[4 ma
	-
Turn over for the next question	



WMP/Jun16/BIOL5

Essay

You should write your essay in continuous prose.

Your essay will be marked for its scientific accuracy. It will also be marked for your selection of relevant material from different parts of the specification and for the quality of your written communication.

The maximum number of marks that can be awarded is

Scientific content	16
Breadth of knowledge	3
Relevance	3
Quality of written communication	3

10 Write an essay on **one** of the following topics.

EITHER

10 (a) The control of processes in cells and the importance of these controls.

[25 marks]

[25 marks]

OR

10 (b) The importance of ions in biology.























END OF QUESTIONS **Copyright Information** For confidentiality purposes, from the November 2015 examination series, acknowledgements of third party copyright material will be published in a separate booklet rather than including them on the examination paper or support materials. This booklet is published after each examination series and is available for free download from www.aqa.org.uk after the live examination series. Permission to reproduce all copyright material has been applied for. In some cases, efforts to contact copyright-holders may have been unsuccessful and AQA will be happy to rectify any omissions of acknowledgements. If you have any queries please contact the Copyright Team, AQA, Stag Hill House,

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