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Centre number	Candidate number
Surname	
Forename(s)	
Candidate signature	5
	I declare this is my own work.

AS BIOLOGY

Paper 2

Time allowed: 1 hour 30 minutes

Materials

For this paper you must have:

- a ruler with millimetre measurements
- · a scientific 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.
- If you need extra space for your answer(s), use the lined pages at the end of this book. Write the question number against your answer(s).
- Show all your working.
- Do all rough work in this book. Cross through any work you do not want to be marked.

Information

- The marks for the questions are shown in brackets.
- The maximum mark for this paper is 75.

For Examiner's Use		
Question	Mark	
1		
2		
3		
4		
5		
6		
7		
8		
9		
TOTAL		

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Answer	all	questions	in	the	spaces	provided

0 1 . 1 The general structure of a fatty acid is RCOOH.

Name the group represented by COOH.

[1 mark]

Carboxyl

Figure 1 shows the structure of a fatty acid R group.

Figure 1

Name the type of R group shown in Figure 1.

Explain your answer.

[2 marks]

Explanation as cit contains a carbon to corbon double

0 1 . 3 Describe how you would test for the presence of a lipid in a liquid sample of food. [2 marks]

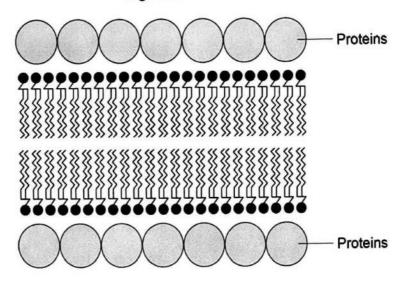
The emulsion test:

add ethanol and then water and shake together. of lipids are present a white 1 milley emulsion is produced.



In 1935, scientists suggested a model for the chemical structure of a cell-surface membrane. **Figure 2** shows the membrane structure the scientists suggested.

Figure 2



0 1 . 4	Give one similarity and two differences between the membrane structure shown in Figure 2 and the fluid-mosaic model of membrane structure. [3 marks]
	Similarity Both have a prospholypid bilayer as its
	port.
	Difference 1 No cholester of like in fluid-mosaic
	model. to intursic protein
	Difference 2 No alycoprotes or glycolipids present like in
	fluid - mosaic model.

Turn over for the next question

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0 2.1	Describe and explain one feature of the alveolar epithelium that makes the epithelium well adapted as a surface for gas exchange. Do not refer to surface area or moisture in your answer. [2 marks]
	It was flattened thin cells and the wall is only one cell thich. This reduces the diffusion
	pathway over which gases have to diffuse across. So the rock of diffusion is much faste,



Doctors measure the health of lungs by calculating the FEV₁:FVC ratio.

FEV₁ is the maximum volume of air exhaled in one second.

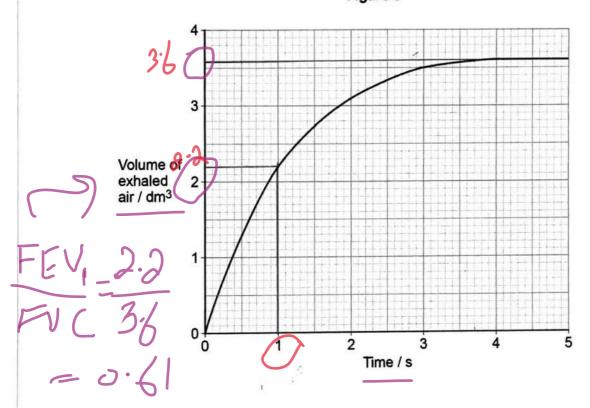
FVC is the maximum volume of air exhaled in one breath.

The minimum FEV₁:FVC ratio of healthy lungs is 0.7:1

Bey meti of eli

A man with the lung disease emphysema inflated his lungs fully. He then exhaled as much of this air as quickly as possible in one breath. Figure 3 shows how the volume of exhaled air changed during this breath.

Figure 3



0 2 2 Use the information provided to determine the FEV₁:FVC ratio of this man's lungs.

Go on to determine how many times greater the minimum ratio of healthy lungs is than

his ratio.

2.3 : 3.6) +3 c

[2 marks]

0.01=1.1475

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FEV₁:FVC ratio of man's lungs = 0.6:1

How many times greater? 1.

1.15

Question 2 continues on the next page



0 2.3

Tidal volume is the volume of air inhaled and exhaled during a single breath when a person is resting. The tidal volume in a person with emphysema is reduced compared with the tidal volume in a healthy person.

Suggest and explain how a reduced tidal volume affects the exchange of carbon dioxide between the blood and the alveoli.

[3 marks]

As there is reduced volume of air eschalad there is more carbon dioxide remaining in the lungs.

This makes the concentration of carbondioxide in the hengs higher, reducing the concentration gradient along which it diffuses out of the blood into the alveoli. This causes slower diffusion rete and more carbon dioxide remaining in the blood.

7



0 3.1	In taxonomy, an organism is identified by referring to the species name and the genus name.
	What term is used to describe this method of naming organisms? [1 mark]
0 3.2	Define the term mutagenic agent. [1 mark]
	A factor that characters the rate at which mutations occur.
0 3 . 3	Figure 4 shows how the species Spartina townsendii is produced.
	The number of chromosomes in cells is shown in some of the boxes.
	Figure 4
	Spartina alterniflora crossed with Spartina maritima Diploid cell 62 Diploid cell
	Gamete 31 Gamete
	Spartina townsendii
	Complete Figure 4 by giving the correct number of chromosomes in each of the boxes. [1 mark]



A mutation in the number of chromosomes in a S. townsendii cell produced a new species, Spartina anglica.

Figure 5 shows the number of chromosomes in leaf cells of these species.

Figure 5

S. townsendii	S. anglica
61 X Z —	122

Name the type of mutation that changed the number of chromosomes in S. townsendii

	to produce S. anglica. Explain your answer.
	Name of mutation Non-disjunction [3 marks]
	Explanation INI Ale share of Mariania in Long Sex Calls
	Explanation At the stage of value see cons
	are being produced chromosomes are not separated
	herraining together in the same cell
0 3.5	Genetic variation within a species is increased during meiosis by crossing over and the independent segregation of homologous chromosomes.
	Apart from mutation, explain one other way genetic variation within a species is
	increased.
	[2 marks]
	Random fertilisation creates a unique/random combination of maternal and paternal
	combination of maternal and paternal
	chronosomes.
	Chariozonics.



0 4.1	Give two structures found in all prokaryotic cells and in all eukaryotic cells. [2 marks]
	1 _ Cyto plasn/ 2 _ Cell surface membrane /
	All prokaryotic cells contain a circular DNA molecule and some prokaryotic cells contain plasmids.
0 4.2	Scientists have found that the rate of plasmid replication is faster in cells growing in a culture with a high concentration of amino acids than in a culture with a lower concentration of amino acids.
	Suggest one explanation for the faster rate of plasmid replication in cells growing in a culture with a high amino acid concentration. [2 marks]
	Armino accids are used for protein synthesis, so with more armino acids more proteins are produced.
	Enzymes involved in DNA replication are proteins, so
	more of these enzymes will lead to faste DNA replication

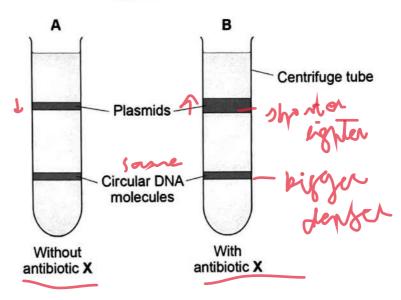


A scientist prepared a culture of a bacterial species.

- She extracted the plasmids and the circular DNA molecules from a sample of cells taken from this culture (A).
- She then added antibiotic X to the culture and let the cells divide for 4 hours.
- She then extracted the plasmids and the circular DNA molecules from a sample of these cells (B).
- The scientist separated the plasmids from the circular DNA molecules in A and in B using ultracentrifugation.

Figure 6 shows her results.

Figure 6



0 4.3	What can you conclude from Figure 6 about a structural difference between the plasmids and the circular DNA? Explain your answer.
	[2 marks]
	Circular DNA is bigger and denser as it's lower
	down the column towards the bottom.
	•



0 4 . 4	replication and on circular DNA replication? Explain your answer. [2 marks]
	As the circular DNA bound is the same width in A
	and B, this surgest its replication stops.
	However for plasmid replication, it continues and
	increases as the band for plasmids is wider
	in B after with X present than without in A.

Turn over for the next question



0 5

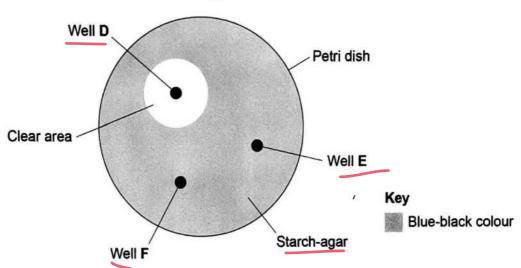
A student investigated the activity of the enzyme amylase. He cut three identical wells (D, E and F) in starch-agar in a Petri dish. He added 0.2 cm³ of:

- amylase solution to well D
- water to well F.

boiled amylase solution to well E

After 60 minutes, he covered the starch-agar with iodine solution. Figure 7 shows his results.

Figure 7



Explain the appearance of the agar in the clear area surrounding well D.

[2 marks]

Amylase hydrolyses starch into more sumple Jodine scheh'an Stains starch blue-black while malkest



0 5.2	What can you conclude about the activity of amylase from the appearance of the agar surrounding well E and well F in Figure 7?
	Uteler is used as a control as there is no anylose
	there is no hydrolysus of starth, so breakdown is
	due to amylase.
	In E, enzyme is clenatured as it has been heated
*	shocker while builting, so it is unable to break down
	Stard.

The student cut out a piece of agar from the clear area surrounding well **D**. He obtained a solution of the substances contained in this piece of agar.

Describe a different biochemical test the student could use with this solution to confirm that amylase had affected the starch in the clear area surrounding well **D**.

[2 marks]

If starch has been broken down by anylase its hydrolysed into maltose, which can be dekated by the Benedicts reagent. Add Benedicts south on to the liquid and heat to at least 80°C. If simple sugars like maltose is present we will see a colow change red/green/orange.

Question 5 continues on the next page

1 5

The diameter of the clear area around well D is 18 mm

In a different investigation, the student prepared a dilution of the amylase solution. He did this by mixing amylase solution and water in the volumes shown in **Table 1**.

Table 1

Amylase solution / cm³	Water / cm³
1.6	2.4

He prepared a starch-agar Petri dish identical to **Figure 7**, but with a single well. He added 0.2 cm³ of the diluted amylase solution to this well and left the Petri dish for 60 minutes.

Use all of this information to predict the diameter of the clear area that will form around the well containing the diluted amylase solution.

Give your answer to the nearest whole number.

Show your working.

1.6 + 2.4 = 4

Dechi of Solution is made

of this 4cm3 1.6cm is anylose

1.6 = 0.4 so 40% of solution is anylose

when 100% is anylose 18mm diameter

 $\times 0.4$ 100% = 18mm 100% = 7.2 100% = 7.2 Answer 100% = 7.2 mm

The student used a ruler to measure the diameter in mm of the clear area around well **D** in **Figure 7**.

Use this information to explain why the answer to Question **05.4** should be given to the nearest whole number.

[1 mark]

[2 marks]

The resolution of a ruler is down to ± 1 mm so

cannot calculate value to higher accuracy.

reparalet

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0 6	The fruit fly is a species of small insect.
	The fruit fly has a gene that codes for an enzyme called alcohol dehydrogenase (AD). AD catalyses the breakdown of alcohol when alcohol is in the insects' food.
	The gene coding for AD has two alleles, ADF and ADS.
06.1	The enzyme encoded by the AD ^s allele catalyses the breakdown of alcohol faster than the enzyme encoded by the AD ^s allele. Suggest why. [3 marks]
	Different genes code for different set primary
	Structure of sequence of aurino acids. Therefore, when
	proteins are folded this differede in primary structure
	will cause different bonds to form, leading to a
	different techning or ever quaterning structure. This
	determines the strape of an enzyme including its
	active site. AD codes for an enzyme that probably
	has a better shape to bid to stabilitate ecusies /faster, so
	more enzure-substrate complexes formed in a given time.
	A scientist took a random sample of adult fruit flies from a population. He measured the frequency of the ADF allele in this sample (generation 0). He then:
	 selected 100 of these insects at random and kept them in a container fed the insects food containing alcohol
	let the insects reproduce
	repeated these steps for 45 generations of fruit fly reproduction.
172	The scientist measured the frequency of the ADF allele in the 45th generation.
0 6.2	Suggest why the scientist took his sample from the population at random. [1 mark]
	Avoid any bias in which individuals
	are sampled.
	Somple will nationally replet exhibited
	y a lopulation



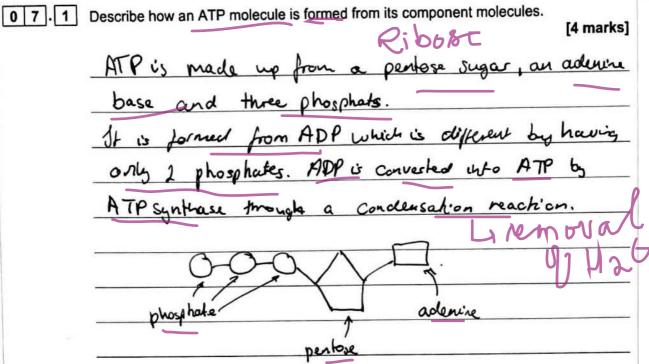
Table 2 shows the scientist's results.

Table 2

Generation of fruit fly reproduction	Frequency of AD
0	0.20
45	0.74

0 6.3	Alcohol is toxic to fruit flies. Suggest and explain why the frequency of the ADF allele changed during the 45 generations. [4 marks]
	Thes with AD7 give can beat down alcohol that is a
	risk to them. This means they have a selective
	advantage over ADS who are less expicient at it.
	This advantage allows ADF gue carrying idividuals
	to toprodu survive beller and reproduce better than
	ADS individuals By doing this they are passing on
	their ADF genes to next generation. Over generations
	this changes the cellele frequency in the population,
	to make the ADF gene more frequent.
0 6.4	Identify the type of selection investigated in the 45 generations of fruit fly reproduction. Tick (✓) one box. [1 mark]
	No selection
	Directional selection Selection forwards getting better at breaking down alcehol.
	Random selection
	Stabilising selection

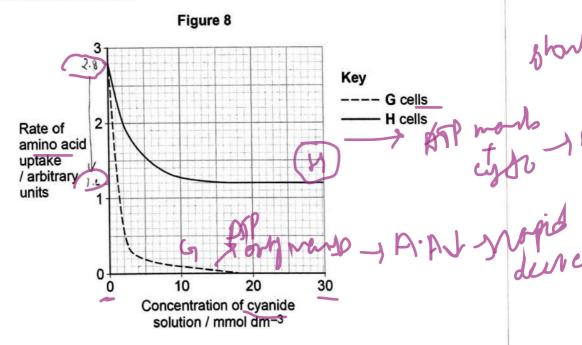




A scientist investigated the effect of cyanide on the rate of amino acid uptake in two types of Escherichia coli, G and H.

- G cells produce enzymes involved in ATP production only on their cell-surface membrane.
- H cells produce enzymes involved in ATP production on their cell-surface membrane and in their cytoplasm.

Figure 8 shows her results.





8

Use Figure 8 to calculate the percentage decrease in the rate of amino acid absorption by H cells in 30 mmol dm⁻³ cyanide solution.

at no cyanicle 2.8 units of amino acid absorbtuon [1 mark] this drops to 1.2 at 30 mmor den-3

 $\frac{2.8 - 1.2 = 1.6}{2.8 \rightarrow 1.6} = \frac{1.6}{2.8} = 0.571428_{\text{Answer}} = \frac{57.1}{96}$ $\Rightarrow \frac{57.1\%}{1.6} = \frac{57.1\%$

Using Figure 8 and the information provided, what can you conclude about amino acid uptake by G cells and by H cells?

Amino acid uptake is done by active transport by

The cell. Cyanode has an effect of reducing and in a color at high enough concentrations stoping amino

acid uptake. As G cells that can only produce

ATP at the membrane Stop taking up amino acids it suggest cyanide stops ATP roduction of ATP at

the membrane which would be needed for achive

transport. But in H cells ATP production can carry

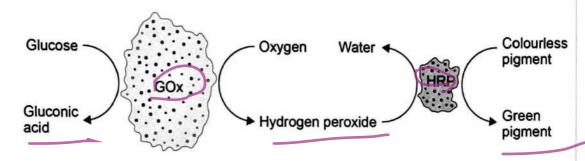
on in the cytoplesm for some mon time.

Turn over for the next question



Figure 9

HRP. Figure 9 shows this sequence of reactions.



A scientist investigated a sequence of reactions catalysed by two enzymes, GOx and

Use **Figure 9** to identify all of the products formed when this sequence of reactions is completed.

[1 mark]

Gluconic acid, green pignent. and water.

The scientist joined DNA molecules together to make tiny cages. The cages are exactly 20 nm long, 20 nm wide and 17 nm deep.

He trapped **one** GOx molecule and **one** HRP molecule together in each cage. The GOx molecule and HRP molecule fill 9% of the cage volume.

The volume of a GOx molecule is eight times larger than an HRP molecule.

Use this information to calculate the volume of a GOx molecule. Give the appropriate unit with your answer.

Show your working.

GOZ +HRP = 9% of cage.

[3 marks]

6

6800 nm x 9 = 612 nm

3612 x 8= 544nm3

Answer 544nm3

1) Volume = 20 × 20 × 17 = 6800 nm3



0 8

The scientist investigated the activity of GOx and HRP enzymes when they are:

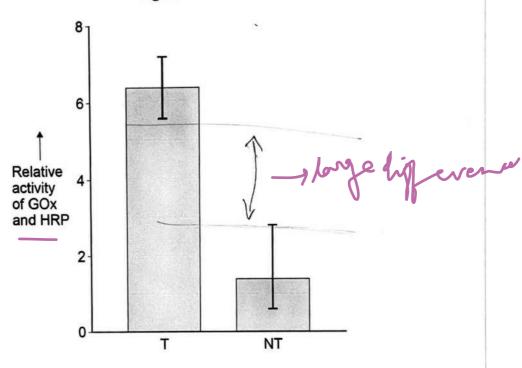
- trapped inside cages (T) and
- not trapped (NT), but free in solution with no cages.

Figure 10 shows his results.

The error bars show ± 2 standard deviations.

± 2 standard deviations include 95% of the data.

Figure 10





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outside the
box

0 8.3	What can you conclude from Figure 10 about the effect of trapping GOx and HRP inside cages? [3 marks]	outside i box
	As the error bars dont overlap we can see there is	
	a significantly higher relative activity when	
	the enzymes are trapped. The error bars represent	
	the Standard alluvations that also about cruckap.	
0 8 . 4	The design of the scientist's investigation did not include a suitable control.	
	Suggest a suitable control. OND [1 mark]	
	A treatment where there is no enzyme actively, so a clenatured form of the enzymes.	
	a denatured form of the enzymes.	
		8

Turn over for the next question

2 5

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0 9 . 1	Explain five properties that make water important for organisms. [5 marks]
	Water is important to all forms of life. This is due
	to its properties like being a great solvent, so metabolic
	reactions can occur it it easier.
	It also has a high heat capacity, this allows organisms
	a buffer from flucturat fluctuations to temperature
	Change.
	Water molecules have a high level of cohesion between
	then. The helps to support water columns in plants,
	which they tell on for saffort structure It also produces
	Surface Lension supporting the structure of small
	organisms as well.
	hastly but not least its an important metabolish itself,
	used in hydrolysis reactions as well as is
	photosynthesis.



0 9 . 2 Describe the process of semi-conservative replication of DNA.

[5 marks]

break

DNA is found in a double helior that needs to have its hydrogen banck broken between base powers before replication. This is done by DNA helicase. Once the two strands are separated each strand can be used to as a temptate to use in semi conservative replication. This nears that on tephicated DNA one Strand of the DNA will be from the Griginal DNA molecule, while the other strand was built by other newcleotroles, based on the original as a temptake. You can use the original strands as a temptate template as nucleotides line up complementary to their base pairs. A pairs with T and Cpairs with G and wise versa. When free nucleotides have lived up confuncting is base pairty to the template strand DNA polymerase joins these needlestides together to form the second strand. These nucleotides are joined by the formation of phosphodiester bonds.

10

END OF QUESTIONS

