

# YEAR 12 UNIT 4

## Topic Test 1 – Heredity

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### Time allowed: 50 minutes Total marks: 40

14 Multiple Choice Questions 4 Short Answer Questions

#### An Answer Sheet is provided for Section A. Answer all questions in Section B in the space provided.

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#### Student Name.....

#### VCE Biology 2014 Year 12 Topic Test 1 Unit 4

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#### **Student Answer Sheet**

There are 14 Multiple Choice questions to be answered by circling the correct letter in the table below. Use only a 2B pencil. If you make a mistake, erase and enter the correct answer. Marks will not be deducted for incorrect answers.

| Question 1  | А | В | С | D | Question 2  | А | В | С | D |
|-------------|---|---|---|---|-------------|---|---|---|---|
| Question 3  | А | В | C | D | Question 4  | A | В | C | D |
| Question 5  | А | В | C | D | Question 6  | A | В | C | D |
| Question 7  | А | В | C | D | Question 8  | A | В | C | D |
| Question 9  | А | В | C | D | Question 10 | A | В | C | D |
| Question 11 | А | В | C | D | Question 12 | A | В | C | D |
| Question 13 | А | В | C | D | Question 14 | A | В | С | D |

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#### **SECTION A – Multiple Choice Questions**

#### **Question 1**

The mRNA transcript of the following DNA sequence - AAATTGC - would be

- A. TTTAACG.
- **B.** GGCCTA.
- C. UUUAACG.
- **D.** TTTUUCG.

Questions 2-4 refer to the diagram shown in Figure 1.



Figure 1

#### **Question 2**

In Figure 1, the structure labelled 2 is the

- A. centriole.
- **B.** nucleus.
- **C.** histone.
- **D.** centromere.

#### **Question 3**

In Figure 1, the structure labelled 1 is a

- A. chromosome.
- **B.** chromatid.
- C. centriole.
- **D.** centromere.

#### **Question 4**

In Figure 1, the most accurate description of the material located in the structure labelled 4 is

- A. chromatin and is composed of DNA wrapped around histone proteins.
- **B.** chromatin and is composed of DNA tightly coiled.
- **C.** a long strand of DNA.
- **D.** chromatin and contains both DNA and RNA wrapped around histone proteins.

*Questions 5 and 6 refer to the following karyotype.* Consider the following karyotype in **Table 1.** 

| χχ | XX | XX      | XX      | XX              | XX             | XX      | XX |
|----|----|---------|---------|-----------------|----------------|---------|----|
|    |    | 3<br>YY | 4<br>YY | 5<br><b>YYY</b> | 6<br><b>YY</b> | 7<br>YY | 8  |
| 9  | 10 | 11      | 12      | 13              | 14             | 15      | 16 |
| XX | XX | XX      | XX      | XX              | XX             | XX      |    |
| 17 | 18 | 19      | 20      | 21              | 22             | 23      | 1  |

Table 1

#### **Question 5**

This karyotype shows a condition termed

- A. trisomy.
- **B.** polyploidy.
- **C.** monosomy.
- **D.** polyteny.

#### **Question 6**

Box 1 contains two chromosomes. How many molecules of DNA does it contain?

- **A.** 2
- **B.** 4
- **C.** 8
- **D.** 1000's

#### **Question 7**

The mitochondrial chromosome is different to the chromosomes in the nucleus because it

- **A.** is single stranded.
- **B.** is circular.
- **C.** contains a greater number of genes.
- **D.** contains sections where there are no genes.

#### **Question 8**

Cystic fibrosis is an autosomal recessive disorder. If a husband and wife are both heterozygous for the affected allele, what is the chance that their firstborn child will have the disease?

- 100% A.
- **B**. 75%
- C. 50%
- D. 25%

#### **Question 9**

A genetic disease is traced back to the paternal grandfather of a family. The mutation causing the disease must have occurred in the

- mitochondrial DNA of a somatic cell. A.
- B. mitochondrial DNA of a germline cell.
- C. nuclear DNA of a somatic cell.
- D. nuclear DNA of a germline cell.

#### **Ouestion 10**

A white, long haired goat (WWLL) is crossed with a black short haired goat (wwll). The offspring are then crossed to produce a third generation. The probability of a white, long haired goat being born in this third generation is

- A.  $\frac{1}{16}$
- **B**.  $\frac{3}{16}$
- C. 9/16
- D.  $\frac{12}{16}$

#### **Question 11**





The mode of inheritance for the trait depicted in the pedigree in Figure 2 is most likely to be

- A. autosomal recessive.
- B. x - linked recessive.
- C. autosomal dominant. D.
- x linked dominant.

#### Question 12 The graph in Figure 3 shows the skin colours of a species of Amazonian frog.



Based on the information in the graph, it would be reasonable to assume that skin colour in this species is a

- A. discontinuous trait controlled by a single gene.
- **B.** discontinuous trait controlled by many genes.
- **C.** continuous trait controlled by a single gene.
- **D.** continuous trait controlled by many genes.

#### **Question 13**

Which of the following scenarios would **not** be an appropriate application for PCR (polymerase chain reaction)?

- **A.** Blood and hair are found at a crime scene and PCR is used to amplify some DNA isolated from cells of these specimens.
- **B.** The karyotype of a person is to be analysed. In preparation, an individual's chromosomes are copied many times by PCR.
- **C.** A tooth of an ancient human is found and a small amount of DNA is obtained. PCR is then used to amplify this fragment for further research.
- **D.** A pig gene is to be inserted into a tomato plant. It is first amplified by PCR before an attempt is made to insert it into tomato cells.

#### **Question 14**

The gene for haemoglobin is exposed to UV light. When the gene is subsequently transcribed and translated only half the haemoglobin protein is manufactured. This suggests that the UV light caused the following type of mutation in the haemoglobin gene.

- A. Translocation.
- **B.** Silent.
- C. Nonsense.
- **D.** Missense.

#### **End of Section A**

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#### **SECTION B – Short Answer Questions**

#### **Question 1 (7 marks)**

In Figure 4, two important cellular processes are illustrated.





| a. | In <b>Figure 4</b> name process A and process B.                                      | 2 marks     |
|----|---|-------------|
| b. | Identify Molecule B in <b>Figure 4</b> and explain why it is shorter than Molecule A. | 2 marks     |
| c. | In <b>Figure 4</b> name Molecule C and explain its function.                          | 2 marks     |
| d. | Identify Molecule D in <b>Figure 4</b> .  | –<br>1 mark |

## Question 2 (7 marks) This question refers to Figure 5 below. Insulin gene Plasmid Bacterium Figure 5



Human insulin can be manufactured by incorporating the human insulin gene into the bacterial cells genome. Some of the components required in this process are illustrated in **Figure 5**.

| What gene           | at tool of genetic engineering must be kept constant, both for the isolation of the e and the cutting of the plasmid?   |
|---------------------|---|
| Ider                | ntify the enzyme that is used to ensure the gene is joined into the plasmid?  |
| Wha                 | at is the term used to describe the process by which bacterial cells take up a foreign e?   |
| Who<br>with<br>succ | en the process is complete there will be many bacteria with the gene and many<br>nout it. Describe the procedure necessary to isolate only the bacteria that have<br>cessfully taken up the gene. |

#### **Question 3 (6 marks)**

The seeds of a species of pea plant have two distinctive traits; they can be round (dominant) or wrinkled (recessive) and yellow (dominant) or green (recessive). If a seed has the phenotype round and yellow, a test cross can be completed to determine its genotype.

#### *Test cross: Yellow, round plant* × *wrinkled, green plant.*

| Why is a test-<br>seeds?       | cross required to determine the genotype of a plant with round, yellow                                 | v<br>1 1 |
|--------------------------------|--|----------|
|                                |  |          |
|                                |  |          |
| Explain how to of the cross an | the test-cross is completed. In your answer, include the possible results nd how they are interpreted. | s 2      |
|                                |  |          |
|                                |  |          |
|                                |  |          |

It is also possible to determine whether the genes are linked or not when a dihybrid cross is completed. The table below shows the genotypes of 100 offspring produced from the following cross:  $GgHh \times gghh$ .

| genotype | Number of offspring |
|----------|---------------------|
| GgHh     | 5                   |
| ggHh     | 45                  |
| Gghh     | 45                  |
| gghh     | 5                   |



\_\_\_\_\_

**d.** Are these two genes linked? Explain your answer.

2 marks

#### **Question 4 (6 marks)**

The pedigree below illustrates the inheritance of a trait in the Spanish Royal family.





a. What is the mode of inheritance of the disease in the pedigree shown in **Figure 6**? Provide a reason for your answer.

2 marks

| What is the genetic status of Eugenie in regard to this disease? Explain.   |        |  |  |  |  |
|---|--------|--|--|--|--|
| If Alfonso marries a woman with no history of the disease in her family and they have a son together, what is the chance that this son will have the disease? | 1 mar  |  |  |  |  |
| Suggest how this disease might have started in this family.   | 1 marl |  |  |  |  |

#### **End of Section B**

End of Topic Test 1

#### **Suggested Answers**

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#### **SECTION A – Multiple Choice Answers**

| 1. C        | <b>2.</b> D | <b>3.</b> B  | <b>4.</b> A  | <b>5.</b> A  | <b>6.</b> B  | <b>7.</b> B  |
|-------------|-------------|--------------|--------------|--------------|--------------|--------------|
| <b>8.</b> D | <b>9.</b> D | <b>10.</b> C | <b>11.</b> A | <b>12.</b> A | <b>13.</b> B | <b>14.</b> C |

#### **SECTION B – Short Answer (Answers)**

#### **Question 1 (7 marks)**

- **a.** Process A is transcription, process B is translation (**2 marks**).
- **b.** Molecule B is mRNA while Molecule A is pre-mRNA (**1 mark**). Molecule A contains introns and exons while Molecule B has had the introns spliced out and only contains exons (**1 mark**).
- **c.** Molecule C is a tRNA molecule (**1 mark**). Its function is to deliver an amino acid to the ribosome in the process of translation (**1 mark**).
- d. Molecule D is a growing polypeptide chain (1 mark).

#### **Question 2 (7 marks)**

- **a.** A plasmid is a small circular piece of double stranded DNA that is contained in bacteria but which lies separate to the bacterial chromosome (**1 mark**). A plasmid can serve as a vector for genes and serves as a means by which genetic material can be incorporated into target cells (**1 mark**).
- **b.** The same restriction enzyme must be used when isolating the gene of interest and cutting open the plasmid prior to inserting the gene (**1 mark**).
- c. DNA ligase (1 mark).
- **d.** Transformations (**1 mark**).
- e. An antibiotic resistance gene is incorporated into the plasmid alongside the gene of interest (1 mark). Thus bacteria can be grown on agar containing the antibiotic and only the bacteria that have successfully taken up the gene will survive and grow (1 mark).

#### Question 3 (6 marks)

- **a.** Round -R, wrinkled -r, Yellow -Y, green -y (**1 mark**).
- **b.** A plant with round, yellow seeds could be homozygous dominant (RRYY) or heterozygous (RrYy) at both these loci. Thus a test cross is required to determine which of these two genotypes is involved (**1 mark**).
- **c.** A plant of the unknown genotype (round, yellow) is crossed with a homozygous recessive plant (wrinkled, green) (**1 mark**). If the resultant offspring are all round and yellow then the genotype in question is homozygous dominant. If some offspring are round and yellow and some wrinkled and green then the genotype of the plant in question is heterozygous at these two loci (**1 mark**).

**d.** Yes, the two genes are linked (**1 mark**). The expected genotypic ratio of this cross is 1:1:1:1. The actual ratio is very different from the expected ratio with two genotypes being heavily overrepresented and the other two underrepresented – this is indicative of linked genes (**1 mark**).

#### **Question 4 (6 marks)**

- **a.** The mode of inheritance of this disease is x-linked recessive (**1 mark**). X-linked because males have the disease much more often than females. This occurs because males only need to inherit one X chromosome bearing a copy of the defective allele to be affected. Recessive because an X chromosome bearing the defective allele can be carried in females when they don't have the disease (**1 mark**).
- **b.** Eugenie is heterozygous for this disease (**1 mark**). She must contain one X chromosome bearing the defective allele as she has sons with the disease who inherit it from her. She also must have one unaffected chromosome as she herself doesn't have the disease (**1 mark**).
- **c.** 0% chance (**1 mark**).
- d. A mutation in a germ line cell or gamete in Beatrice's parent or ancestor. (1 mark).

#### **End of Suggested Answers**