



Additional Assessment Materials  
Summer 2021

Pearson Edexcel GCSE in Biology (1BI0)  
Higher

Resource Set Topic 3: Genetics

Questions

(Public release version)

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## General guidance to Additional Assessment Materials for use in 2021

### Context

- Additional Assessment Materials are being produced for GCSE, AS and A levels (with the exception of Art and Design).
- The Additional Assessment Materials presented in this booklet are an **optional** part of the range of evidence teachers may use when deciding on a candidate's grade.
- 2021 Additional Assessment Materials have been drawn from previous examination materials, namely past papers.
- Additional Assessment Materials have come from past papers both published (those materials available publicly) and unpublished (those currently under padlock to our centres) presented in a different format to allow teachers to adapt them for use with candidate.

### Purpose

- The purpose of this resource to provide qualification-specific sets/groups of questions covering the knowledge, skills and understanding relevant to this Pearson qualification.
- This document should be used in conjunction with the mapping guidance which will map content and/or skills covered within each set of questions.
- These materials are only intended to support the summer 2021 series.

7.

(b) Mendel's research on pea plants showed that genetic traits are inherited.

(i) Which term is used to describe the expression of traits in an organism?

(1)

- A** genotype
- B** phenotype
- C** allele
- D** gamete

(ii) Mendel crossed pea plants that produced round seeds with pea plants that produced wrinkled seeds.

All the offspring produced round seeds.

He then crossed these offspring with each other.

Some pea plants in the next generation produced round seeds and the others produced wrinkled seeds.

Explain how this showed that some inherited traits are not expressed in an organism.

(3)

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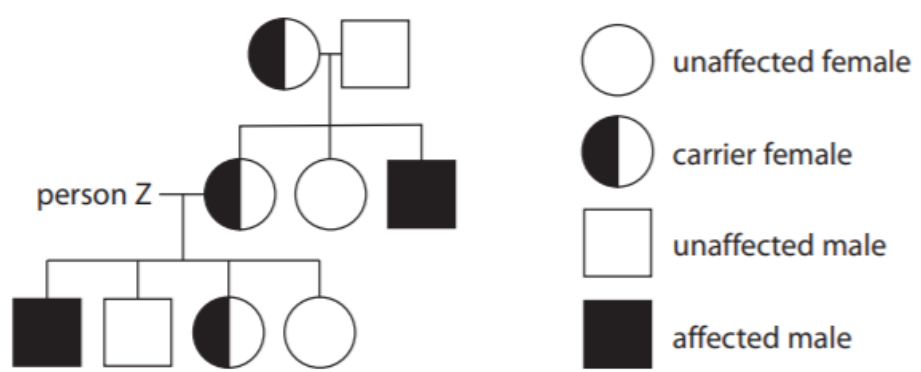
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7.

(c) Duchenne muscular dystrophy is a recessive sex-linked genetic disorder. This disorder causes muscle weakness.

Figure 14 shows the inheritance of Duchenne muscular dystrophy in a family.



**Figure 14**

State and explain the phenotype of person Z.

(3)

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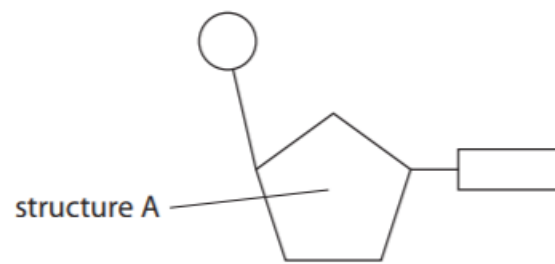
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(b) Figure 17 shows the structure of a DNA nucleotide.



**Figure 17**

(i) Structure A is a

- A** base
- B** phosphate
- C** sugar
- D** polymer

(1)



(ii) Myxopyronin inhibits bacterial RNA polymerase.

Explain why the antibiotic myxopyronin can be used to treat bacterial infections in humans.

(4)

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**2** (a) A scientist obtained a mass of 0.0062 nanograms of DNA from a diploid human cell.

Calculate the mass of DNA the scientist should obtain from a haploid human cell.

Give your answer in picograms.

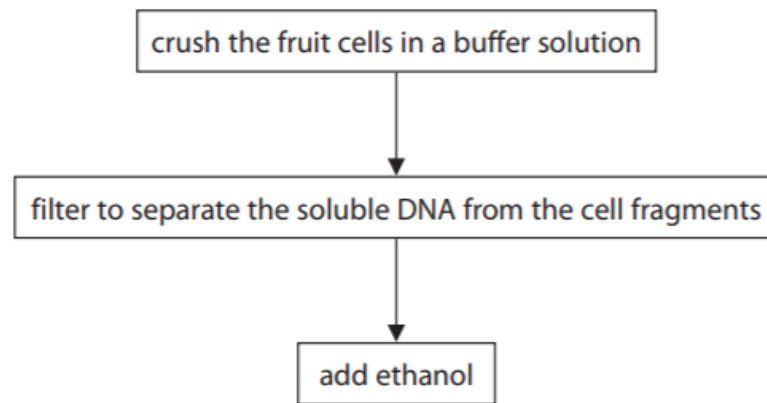
(1 nanogram = 1000 picograms)

(2)

..... picograms



(b) A student used the method shown in Figure 3 to compare the mass of DNA extracted from strawberry fruit cells and from kiwi fruit cells.



**Figure 3**

(i) State why ethanol is used.

(1)

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(ii) State **two** variables the student needs to control when using this method to compare the mass of DNA from these two fruits.

(2)

1 \_\_\_\_\_

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2 \_\_\_\_\_

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(iii) The student repeated the experiment.

Give **one** reason why.

(1)

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(c) Mitosis and meiosis are processes that produce new cells.

Compare the outcomes of mitosis and meiosis.

(3)

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9 Transcription and translation are stages in the synthesis of proteins.

(a) (i) Which enzyme is involved in the process of transcription?

(1)

- A DNA ligase
- B lysozyme
- C RNA polymerase
- D restriction endonuclease

(ii) Describe how a mutation in the non-coding region of the DNA can prevent a gene being transcribed.

(2)

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(b) A gene coding for a protein has two alleles.

Figure 14 shows the first 5 codons of an mRNA strand for these alleles.

Allele 1 – AUG CCA CAG GAG UUC

Allele 2 – AUG CCA GAG GAG UUC

**Figure 14**

Allele 2 has a mutation.

Figure 15 shows the key needed to predict the translated amino acid sequence of the protein.

<b>codon</b>	AUG	CCA	CAG	GAG	UUC
<b>amino acid</b>	Met	Pro	Gln	Glu	Phe

**Figure 15**

Explain how the mutation in allele 2 could affect the functioning of this protein.

(3)

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\*(c) The inheritance of different alleles affects the phenotype of an individual.

A child is blood group O.

The child's mother is blood group A and the child's father is blood group B.

Explain how this child is blood group O.

Use the Punnett square and probability in your answer.

(6)


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(ii) State what is meant by the term genome.

(1)

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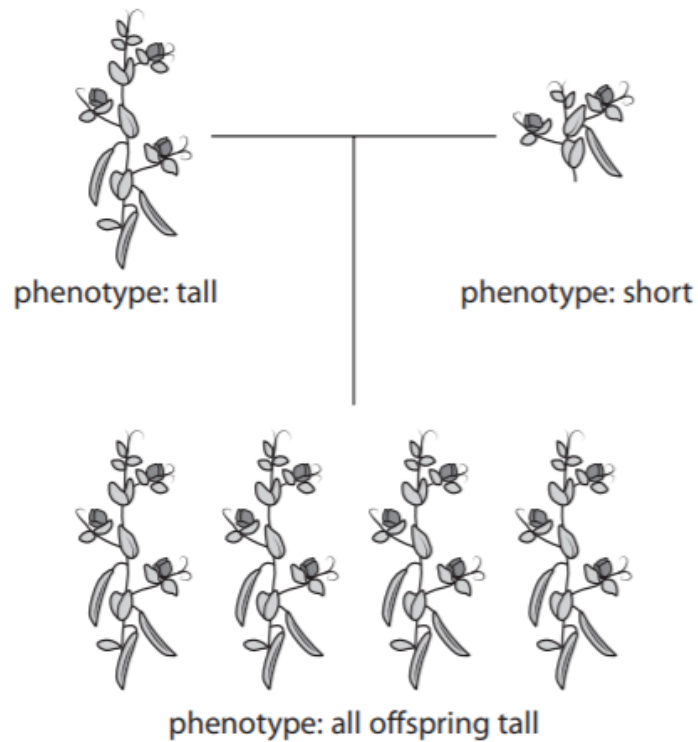
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4 Gregor Mendel used pea plants in plant breeding experiments. He discovered the basis of genetic inheritance.

(a) He cross-bred tall pea plants with short pea plants.

All the offspring were tall, as shown in Figure 4.



**Figure 4**

(i) Explain why the offspring are all tall.

(2)

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(ii) In this investigation, the parent pea plants were grown in a warm, closed greenhouse.

Give **two** reasons why the parent pea plants were grown in a warm, closed greenhouse.

(2)

1 .....

2 .....

(b) Pea plants produce different coloured peas.

The allele for yellow-coloured peas (A) is dominant to the allele for green-coloured peas (a).

Two heterozygous parent plants were used in a genetic cross.

(i) Predict, using the Punnett square, the percentage probability that this cross will have offspring that produce green-coloured peas.

(3)


percentage probability of green-coloured peas = .....%



(ii) Which statement correctly describes a DNA molecule?

(1)

- A** two strands joined together by strong bonds to form a double helix
- B** two complementary bases twisted into a double helix by strong bonds
- C** a double helix with strands joined by hydrogen bonds between bases
- D** four complementary strands joined together with hydrogen bonds

(iii) State the term used to describe a change in the sequence of DNA bases.

(1)

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10.

(b) Colour blindness affects approximately 1 in 12 men.

In a city of 2 million people, 51% are men.

(i) What is the number of men who are colour blind in the city?

(1)

- A** 42 500
- B** 85 000
- C** 166 666
- D** 1 020 000

(ii) Colour blindness is a sex-linked genetic disorder caused by a recessive allele.

Colour blindness only affects 1 in 200 women.

Explain why more men than women are colour blind.

(2)

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(iii) A female without the allele for colour blindness has a baby boy.

The father is colour blind.

Explain the probability of the baby boy being colour blind.

(2)

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10.

(c) One cause of colour blindness is a change in the DNA sequence of a gene.

This results in the production of a different protein in cone cells in the retina of the eye.

Explain how a change in the DNA sequence of a gene can result in the production of a different protein.

(4)

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**TOTAL = 61 MARKS**