



Additional Assessment Materials
Summer 2021

Pearson Edexcel GCE in A Level Biology

Topic 8: Origins of Genetic Variation

(Public release version)

Pearson: helping people progress, everywhere

Pearson aspires to be the world's leading learning company. Our aim is to help everyone progress in their lives through education. We believe in every kind of learning, for all kinds of people, wherever they are in the world. We've been involved in education for over 150 years, and by working across 70 countries, in 100 languages, we have built an international reputation for our commitment to high standards and raising achievement through innovation in education. Find out more about how we can help you and your students at: www.pearson.com/uk

General guidance to Additional Assessment Materials for use in 2021

Context

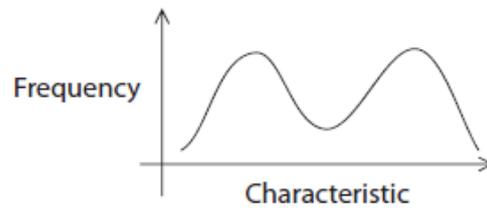
- Additional Assessment Materials are being produced for GCSE, AS and A levels (with the exception of Art and Design).
- The Additional Assessment Materials presented in this booklet are an **optional** part of the range of evidence teachers may use when deciding on a candidate's grade.
- 2021 Additional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- Additional Assessment Materials have come from past papers both published (those materials available publicly) and unpublished (those currently under padlock to our centres) presented in a different format to allow teachers to adapt them for use with candidate.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

1

(a) The graph shows the frequency of a characteristic found in a population of animals.



Which type of selection would create this pattern?

(1)

- A allopatric selection
- B directional selection
- C disruptive selection
- D stabilising selection

(b) The Eurasian lynx is the largest native European cat species.

It was once widespread across Europe but is now restricted to small areas of national parks.



Ex-situ and *in-situ* conservation measures were used in the 1970s to increase biodiversity.

- Lynx were bred in zoos and 10 were reintroduced into an area of protected forest where the lynx had become extinct.
- Existing wild lynx were protected in an area of forest where they had not become extinct.

In 2016 scientists estimated the population sizes and genetic biodiversity of the lynx in these two areas of forest.

They found that the population and genetic biodiversity of the lynx in the area where they had been reintroduced were much lower than in the protected area.

(i) State what is meant by the term **biodiversity**.

(1)

The number and variety of living organisms in a given region.

(ii) Explain why, in 2016, the genetic biodiversity of the lynx population in the area where they had been reintroduced was much lower than in the protected area.

(2)

Only a small number of lynx were reintroduced into the area of protected forest. These were bred in a zoo in which there would be a limited gene pool and inbreeding would likely occur, resulting in low genetic biodiversity.

(c) Explain the principles and issues associated with *ex-situ* conservation methods.

(4)

Ex-situ conservation methods involve the conservation of all biodiversity levels outside their natural habitats. Zoos, botanical gardens and seed banks are aimed at conserving endangered species and their genetic diversity, whilst captive breeding programmes focus on increasing population numbers. Such methods also involve educating the public about the need for conservation. However, there are ethical issues associated with ex-situ methods. For example, changes in animal behaviour (e.g. increased aggression, self-mutilation) have been observed in zoos. Deterioration of genetic diversity and inbreeding is also a problem although this can be overcome using methods such as artificial insemination.

2

In the fruit fly, *Drosophila*, the allele for normal wings (N) is dominant to the allele for vestigial (small) wings (n).

The allele for red eyes (R) is dominant to the allele for sepia eyes (r).

In an investigation, students crossed homozygous parent flies. Flies with normal wings and red eyes were crossed with flies with vestigial wings and sepia eyes.

All the F₁ offspring of this cross had normal wings and red eyes.

Flies from this F₁ generation were crossed and the phenotypes of their offspring (F₂ generation) were counted.

The results for the F₂ generation are shown in the table.

| <i>Drosophila</i> phenotype | Number of <i>Drosophila</i> with each phenotype |
|--------------------------------|---|
| normal wings and red eyes | 885 |
| normal wings and sepia eyes | 322 |
| vestigial wings and red eyes | 286 |
| vestigial wings and sepia eyes | 107 |

The students thought that the genes for wing length and eye colour were on different chromosomes.

(a) (i) State a null hypothesis for this investigation.

(1)

No significant difference between observed and expected numbers of offspring phenotypes.

(ii) A Chi squared test was carried out to test this hypothesis.

Complete the table.

(1)

| Phenotype | Expected ratio | Observed results (O) | Expected results (E) | (O - E) | (O - E) ² | $\frac{(O - E)^2}{E}$ |
|--------------------------------|----------------|----------------------|----------------------|---------|----------------------|-----------------------|
| normal wings and red eyes | 9 | 885 | 900 | -15 | 225 | 0.25 |
| normal wings and sepia eyes | 3 | 322 | 300 | 22 | 484 | 1.61 |
| vestigial wings and red eyes | 3 | 286 | 300 | -14 | 196 | 0.65 |
| vestigial wings and sepia eyes | 1 | 107 | 100 | 7 | 49 | 0.49 |

(iii) Calculate the value of Chi squared using the formula

(1)

$$\chi^2 = \sum \frac{(O - E)^2}{E}$$

$$\chi^2 = 0.25 + 1.61 + 0.65 + 0.49 = 3.00$$

Answer 3

(iv) The table shows some critical values of Chi squared at different degrees of freedom.

| Degrees of freedom | p value | | | | |
|--------------------|---------|-------|-------|-------|--------|
| | 0.900 | 0.500 | 0.100 | 0.050 | 0.010 |
| 1 | 0.016 | 0.455 | 2.706 | 3.841 | 6.635 |
| 2 | 0.211 | 1.386 | 4.605 | 5.991 | 9.210 |
| 3 | 0.584 | 2.366 | 6.251 | 7.815 | 11.345 |
| 4 | 1.064 | 3.357 | 7.779 | 9.488 | 13.277 |

Use this table to comment on the results of the investigation.

(3)

Probability of 0.05, df of 3, critical value is 7.815. $3 < 7.815$.
H₀ is accepted. There is no significant difference between
observed and expected numbers of offspring phenotypes.

*(b) In *Drosophila*, the allele for grey bodies (G) is dominant to the allele for black bodies (g).

In a second investigation, students crossed homozygous parent flies. Flies with normal wings and grey bodies were crossed with flies with vestigial wings and black bodies.

All the F₁ offspring had normal wings and grey bodies.

Flies from this F₁ generation were crossed and the phenotypes of their offspring (F₂ generation) were counted.

The results are shown in the table.

| <i>Drosophila</i> phenotype | Number of <i>Drosophila</i> with each phenotype |
|--------------------------------|---|
| normal wings and grey body | 1105 |
| normal wings and black body | 85 |
| vestigial wings and grey body | 72 |
| vestigial wings and black body | 338 |

Explain the results of this second investigation.

(6)

Parent genotypes are NNGG and nngg. The F₁ generation are therefore all heterozygous, NnGg, with normal wings and grey bodies. A ratio of 9:3:3:1 would be expected in the F₂ generation. However, this was not observed, with the number of parental phenotypes greater than recombinant phenotypes. This is likely due to autosomal gene linkage, with genes for body colour and wing size located close together on the same autosome. They are unlikely to undergo recombination during meiosis so are inherited together. Members of the F₁ generation have one chromosome with GN and one with gn so most gametes will have GN or gn allele combinations. A low frequency of recombinant gametes are produced as a result of crossing over during prophase I of meiosis in which DNA is exchanged between homologous chromosomes at chiasmata.

This produces a small number of gametes with Ng and ng allele combinations. These are responsible for the small number of normal wing, black bodied and vestigial wing, grey bodied *Drosophila*. A greater number of normal wing, grey bodied *Drosophila* are observed than vestigial wing, black bodied due to the dominant GN alleles which override gn.

3

The photograph shows a maize cob with smooth, wrinkled and different coloured grains.



© W.P. Armstrong 2001

The shape and colour of maize grains are controlled by two unlinked genes.

The allele for smooth seeds (A) is dominant to the allele for wrinkled seeds (a).

The allele for purple seeds (B) is dominant to the allele for yellow seeds (b).

(a) State all the possible genotypes of a smooth, purple grain.

(1)

AABB, AaBB, AABb, AaBb

(b) Two maize plants, grown from grains that were both wrinkled and purple, were cross-pollinated.

In the F_1 generation, some grains were wrinkled and purple and some were wrinkled and yellow.

Which of the following shows the genotypes of the parent plants?

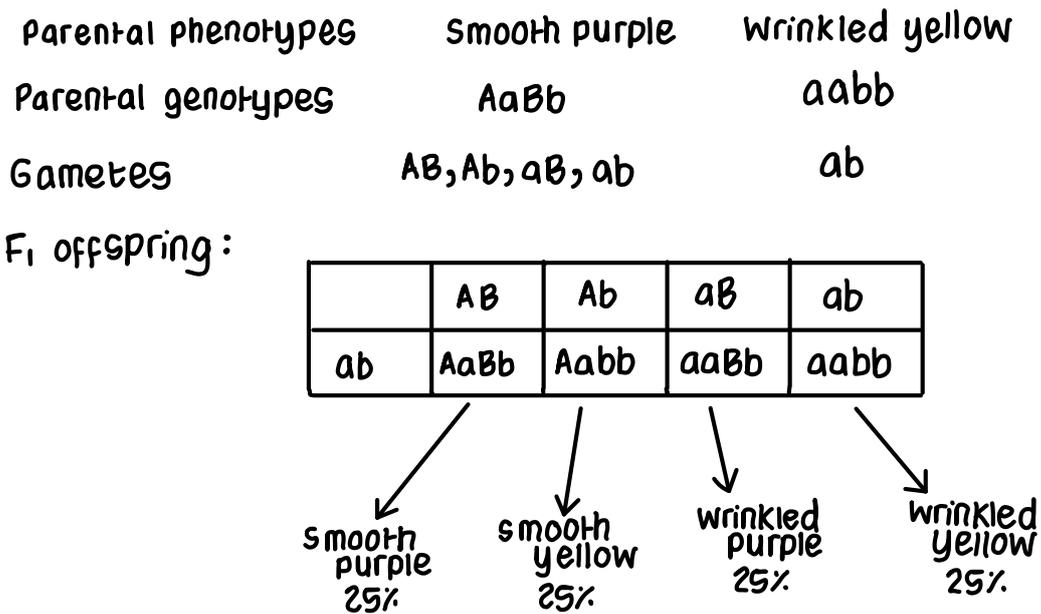
(1)

- A $AaBb \times AaBb$
- B $aaBB \times aaBB$
- C $AABb \times AABB$
- D $aaBb \times aaBb$

(c) A student cross-pollinated a maize plant grown from a smooth, purple grain (heterozygous for both pairs of alleles) with a maize plant grown from a wrinkled, yellow grain.

Using a genetic diagram, determine the probability that this cross will produce grains that are wrinkled and purple.

(4)



Answer 25%

4

Cystic fibrosis is a recessive inherited condition where the cells in the lungs produce sticky mucus. This mucus builds up in the airways, causing breathlessness and chest infections.

People with cystic fibrosis often need treatments such as physiotherapy and antibiotics.

(a) The incidence of babies born with cystic fibrosis in Australia is 1 in 2500.

Use the Hardy Weinberg equation, $p^2 + 2pq + q^2 = 1$, to calculate the percentage of Australians who are carriers of cystic fibrosis.

(4)

$$q^2 = \frac{1}{2500} = 0.0004$$

$$q = \sqrt{0.0004} = 0.02$$

$$p = 1 - q = 1 - 0.02 = 0.98$$

$$2pq = 2 \times 0.02 \times 0.98 = 0.0392$$

$$0.0392 \times 100 = 3.92\%$$

Answer 3.92 %

(b) A woman is a carrier of the cystic fibrosis allele. Her partner does not have cystic fibrosis and is not a carrier.

Use a genetic cross to determine the probability of this woman producing a child who is also a carrier.

Parental genotypes ♀ ♂ (4)

 Cc CC

Gametes (C) (c) (C) (C)

F₁ genotypes :

| | | |
|---|----|----|
| | C | C |
| C | CC | CC |
| c | Cc | Cc |

∴ 50% Cc carriers
50% CC normal

Probability 50%

Some health disorders, such as sickle cell anaemia, have a genetic basis.

People who are at risk of these disorders can be identified using genetic tests.

Hospital managers need to predict the future cost of treating people with health disorders.

It has been claimed that the Hardy-Weinberg equation ($p + q = 1$ or $p^2 + 2pq + q^2 = 1$) could be used to predict the number of people who would need treatment for health disorders.

Discuss the validity of this claim.

(9)

The Hardy-Weinberg equation assumes that the frequency of alleles remains constant, $p + q = 1$, where p is the frequency of the dominant allele and q is the frequency of the recessive allele. It also assumes that the frequency of genotypes stays constant, $p^2 + 2pq + q^2 = 1$, where p^2 is the frequency of homozygous dominant, $2pq$ is the frequency of heterozygous and q^2 is the frequency of homozygous recessive. With the knowledge of dominant and recessive alleles, hospital managers could predict the number of people who may need treatment for purely genetic disorders. However, not all of the principle's assumptions will be met. The HW principle assumes no new mutations, no immigration or emigration, no natural selection for or against alleles, a large population and random mating.

Assumptions such as no migration will not hold in the majority of human populations. Moreover, populations differ in size, and in small countries for example, genetic drift will have an effect. Thus, use of the HW principle is not valid. Additionally, although sickle cell anaemia has a genetic basis, other health disorders such as heart disease and various types of cancers do not solely depend on genetics. Other lifestyle factors may be involved e.g. smoking increases the risk of lung cancer. Thus the HW principle cannot

