

Surname
First name(s)

Date

Tutor



GCE A LEVEL

BW-A2-4.3-W1

TOPIC WORKSHEET 1 – 4.3 INHERITANCE

BIOLOGY – A2 unit 4

Variation, Inheritance and Options

Suggested time: 3 hours 15 minutes

For Tutor's use only

Question	Maximum Mark	Mark Awarded	Question	Maximum Mark	Mark Awarded
1.	9		9.	10	
2.	10		10.	9	
3.	10		11.	10	
4.	9		12.	9	
5.	10		13.	11	
6.	10		14.	9	
7.	11		15.	9	
8.	9		Total	145	

ADDITIONAL MATERIALS

In addition to this worksheet, you will require a calculator and a ruler.

INSTRUCTIONS TO CANDIDATES

Use black ink or black ball-point pen. Do not use gel pen or correction fluid.

You may use a pencil for graphs and diagrams only.

Write your name and the date in the spaces at the top of this page.

Answer **all** questions.

Write your answers in the spaces provided in this booklet. If you are working on screen, write your answers on paper and number each one clearly.

INFORMATION FOR CANDIDATES

The number of marks is given in brackets at the end of each question or part-question. The worksheet is out of **145** marks.

The assessment of the quality of extended response (QER) will take place in question **15**.

The quality of written communication will affect the awarding of marks.

This worksheet is written by BioWelsh in the style of the WJEC A level Biology examination, following the WJEC specification (topic 4.3). It is not an official WJEC paper.



Answer **all** questions.

1. In the fruit fly *Drosophila melanogaster*, a gene for body colour has two alleles. The allele for a grey body, **E**, is dominant to the allele for a black (ebony) body, **e**. **Image 1.1** shows a pair of homologous chromosomes from a body cell of one fly, with the alleles of three of its genes shown on the chromosomes.

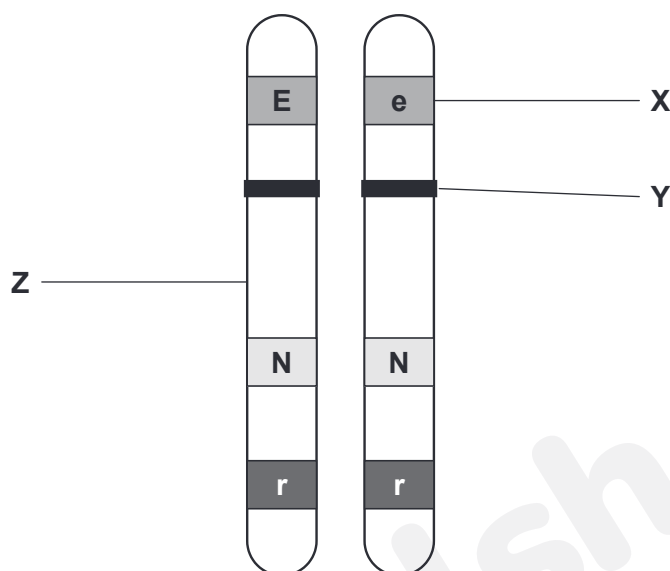


Image 1.1

- (a) Identify the structures or regions labelled **X**, **Y** and **Z**. [3]

X
Y
Z

- (b) Define the term **allele**. [1]

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- (c) Complete **Table 1.2** to show the genotype of this fly for each gene, and whether it is homozygous or heterozygous for that gene. [2]

Table 1.2

Gene	Genotype	Homozygous or heterozygous
body colour		
gene N		
gene R		

- (d) This fly is crossed with a fly that has a black body. Draw a genetic diagram to show the genotypes and phenotypes of the offspring you would expect, and state the expected phenotype ratio. [3]

2. In Andalusian chickens, feather colour is controlled by one gene with two codominant alleles, C^B and C^W . Birds that are $C^B C^B$ have black feathers, birds that are $C^W C^W$ are white with a few dark flecks (called splash), and birds with both alleles have blue-grey feathers and are called blue. A breeder mated blue birds with each other. **Table 2.1** shows the chicks that hatched.

Table 2.1

Feather colour of chicks	Number of chicks
black	23
blue	51
splash	26

- (a) Explain what is meant by codominant alleles, using the blue chickens as an example. [2]

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- (b) Draw a genetic diagram to show why the breeder obtained the results in **Table 2.1**. [3]

- (c) Use **Table 2.1** to state the ratio of black to blue to splash chicks, as a whole-number ratio. [1]

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- (d) Explain why the breeder can never obtain a flock in which all the offspring of blue parents are blue. [2]

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- (e) Suggest which birds the breeder should mate to obtain only blue chicks. Explain your answer. [2]

3. Manx cats have no tail. Taillessness is caused by a dominant allele, **M**. Embryos that are homozygous **MM** die early in development, so no **MM** kittens are born. Cats that are **mm** have a normal tail. **Table 3.1** shows the total number of kittens born in several litters from matings between two Manx cats.

Table 3.1

Phenotype of kittens	Number of kittens
tailless (Manx)	58
tailed	31
total	89

- (a) State the genotype of every Manx cat that is born. Give a reason for your answer. [1]

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- (b) Draw a genetic diagram to predict the phenotype ratio of kittens born when two Manx cats are mated. [3]

A student wrongly expected a ratio of 3 tailless : 1 tailed and used a chi-squared test on the results in **Table 3.1**, using the formula:

$$\chi^2 = \sum (O - E)^2 \div E$$

- (c) (i) Calculate the value of χ^2 for the student's hypothesis. Show your working. [3]

$$\chi^2 = \dots\dots\dots$$

- (ii) The critical value of χ^2 for 1 degree of freedom at $p = 0.05$ is 3.84. State **and** explain what the value of χ^2 shows about the student's hypothesis. [2]

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- (iii) Explain how the results in **Table 3.1** support the idea that **MM** embryos die. [1]

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4. **Image 4.1** shows a cell from an animal during meiosis. The cell has the genotype AaBb, and the two genes are on different chromosomes.

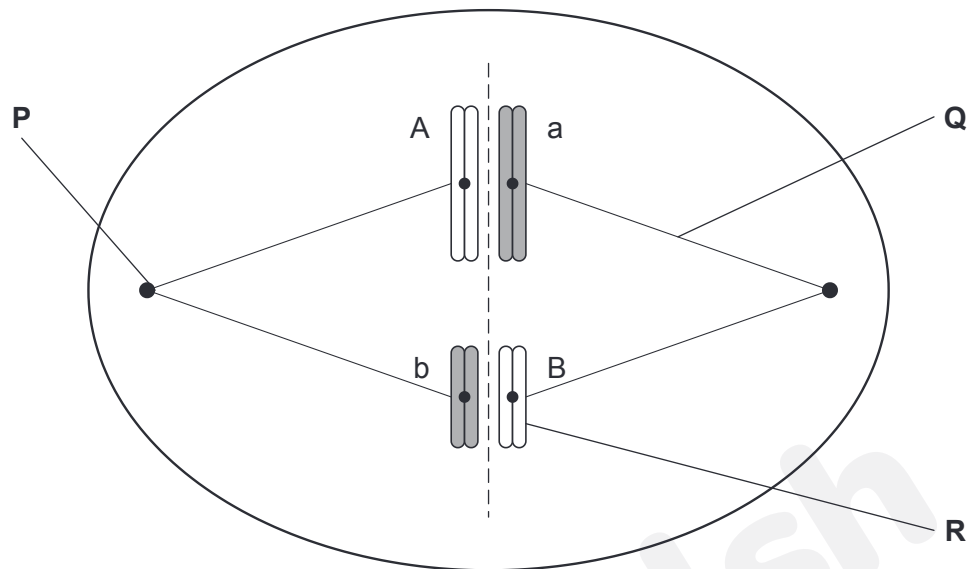


Image 4.1

- (a) Name the stage of meiosis shown in **Image 4.1**. Give a reason for your answer. [2]

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- (b) Identify the structures labelled **P**, **Q** and **R**. [2]

P

Q

R

- (c) State the genotypes of the gametes that will be produced from the arrangement shown in **Image 4.1**. [1]

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- (d) Explain how this animal is able to produce four genetically different types of gamete in equal numbers. [3]

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- (e) The body cells of this animal have 8 chromosomes. Calculate the number of different combinations of chromosomes that are possible in its gametes as a result of independent assortment alone. [1]

Number of combinations =

5. In *Drosophila melanogaster*, a grey body (**E**) is dominant to an ebony body (**e**), and normal wings (**V**) are dominant to vestigial wings (**v**). The two genes are on different chromosomes. Flies that were heterozygous for both genes were crossed with each other. **Table 5.1** shows the offspring.

Table 5.1

Phenotype	Number of flies
grey body, normal wings	290
grey body, vestigial wings	101
ebony body, normal wings	92
ebony body, vestigial wings	37
total	520

- (a) State the genotypes of the gametes produced by one of the heterozygous parent flies. [1]

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- (b) State the phenotype ratio expected in the offspring, and explain why this ratio is expected. [2]

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The null hypothesis was that there is no significant difference between the observed numbers and the numbers expected from this ratio.

- (c) (i) Complete **Table 5.2** and calculate χ^2 . [3]

Table 5.2

Phenotype	Observed (O)	Expected (E)	$(O - E)^2 \div E$
grey, normal	290	292.5	0.02
grey, vestigial	101		0.13
ebony, normal	92	97.5	
ebony, vestigial	37		
		$\chi^2 =$	

Table 5.3 shows critical values of χ^2 at $p = 0.05$.

Table 5.3

Degrees of freedom	1	2	3	4
Critical value	3.84	5.99	7.82	9.49

- (ii) State the number of degrees of freedom, and use **Table 5.3** to explain whether the null hypothesis should be accepted or rejected. [3]

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- (d) Explain why the test is more reliable when a large number of flies is counted. [1]

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6. In sweet peas (*Lathyrus odoratus*), purple flowers (**P**) are dominant to red flowers (**p**), and long pollen grains (**L**) are dominant to round pollen grains (**l**). Early in the twentieth century, geneticists crossed plants that were heterozygous for both genes, and obtained the offspring shown in **Table 6.1**.

Table 6.1

Phenotype	Number of plants
purple flowers, long pollen	4831
purple flowers, round pollen	390
red flowers, long pollen	393
red flowers, round pollen	1338
total	6952

- (a) Calculate the number of plants with purple flowers and long pollen that would be expected if the ratio were 9 : 3 : 3 : 1. [1]

Expected number =

- (b) The value of χ^2 for these results, tested against 9 : 3 : 3 : 1, is 3372. The critical value at $p = 0.05$ is 7.82. State the number of degrees of freedom and what this value of χ^2 shows. [3]

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- (c) Explain why many more plants had purple flowers with long pollen, and red flowers with round pollen, than a 9 : 3 : 3 : 1 ratio predicts. [3]

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- (d) Explain how the plants with purple flowers and round pollen, and with red flowers and long pollen, were produced. [2]

- (e) Suggest what the small number of these plants shows about the positions of the two genes on the chromosome. [1]

7. Tobacco seeds were collected from a plant that was heterozygous for a gene needed to make chlorophyll. The allele for chlorophyll production, **G**, is dominant to the allele **g**; seedlings that are **gg** are albino (white). The plant had been self-pollinated. Students placed 55 seeds on damp filter paper in each of five Petri dishes, kept them in the light at 25 °C and counted the seedlings after 10 days. **Table 7.1** shows their results.

Table 7.1

Dish	1	2	3	4	5	Total
Green seedlings	38	41	36	40	37	192
Albino seedlings	14	12	15	11	16	68

- (a) Explain why the dishes were kept in the light. [2]

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- (b) Suggest **two** variables, other than light and temperature, that should be kept the same in each dish. [2]

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The students used this formula:

$$\chi^2 = \sum (O - E)^2 \div E$$

- (c) Use the totals in **Table 7.1** to calculate χ^2 for a ratio of 3 green : 1 albino. [3]

$$\chi^2 = \dots\dots\dots$$

- (d) The critical value of χ^2 for 1 degree of freedom at $p = 0.05$ is 3.84. State the conclusion the students should make. [2]

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- (e) The students counted the seedlings again after three weeks and found far fewer albino seedlings. Explain why. [2]

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8. **Image 8.1** shows the chromosomes from a body cell of a man, photographed and arranged in pairs.



Image 8.1

- (a) State the number of autosomes in this cell. [1]

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- (b) Identify chromosomes **T** and **U**, and explain how they show that the cell came from a male. [2]

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- (c) Explain why the father, and not the mother, determines the sex of a child. [2]

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Red-green colour blindness is caused by a recessive allele, X^b , of a gene on the X chromosome. The normal allele is X^B . A woman with normal vision, whose father was colour blind, has children with a man with normal vision.

- (d) (i) Explain why the woman must be a carrier. [1]

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- (ii) Draw a genetic diagram to find the probability that one of their sons is colour blind. [3]

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9. Haemophilia A is caused by a recessive allele of a gene for a blood clotting protein on the X chromosome. **Image 9.1** shows the inheritance of haemophilia A in one family.

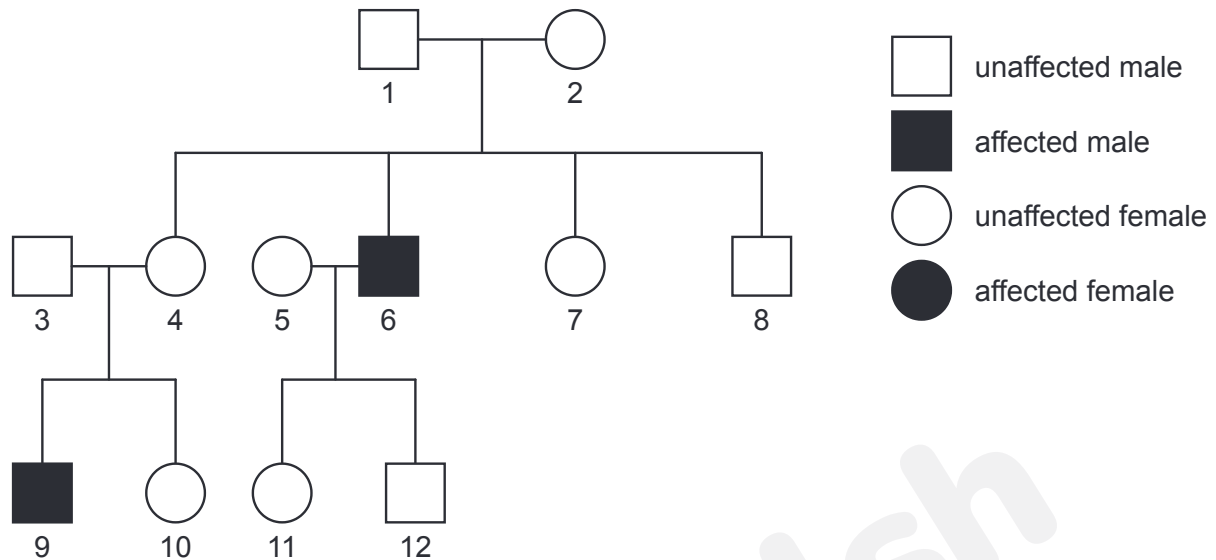


Image 9.1

- (a) Explain why haemophilia A is much more common in males than in females. [2]

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- (b) Using X^H for the normal allele and X^h for the haemophilia allele, give the genotypes of individuals **2**, **6** and **11**. [3]

2

6

11

- (c) Explain why individual **4** must be a carrier even though she does not have haemophilia. [2]

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- (d) Individual **11** has a child with a man who does not have haemophilia. Calculate the probability that the child is a boy with haemophilia. [1]

Probability =

- (e) Explain why the genotype of individual **7** cannot be determined from **Image 9.1**. [2]

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10. Some forms of inherited deafness are caused by a single gene. **Image 10.1** shows a family in which one form of inherited deafness occurs.

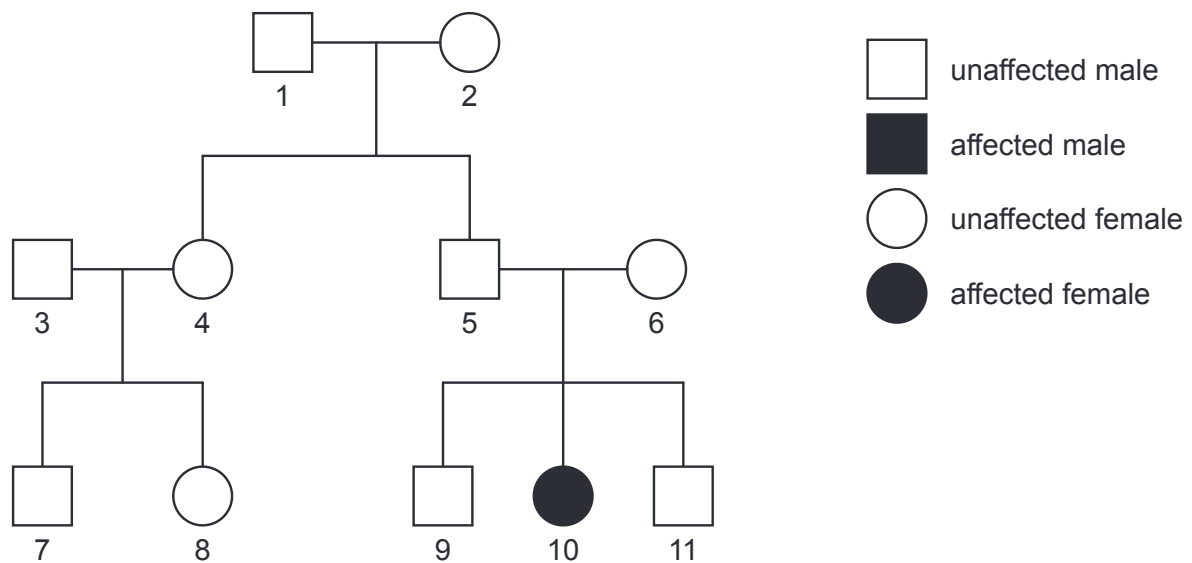


Image 10.1

- (a) Using evidence from **Image 10.1**, explain why the allele causing this form of deafness is recessive. [2]

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- (b) Using evidence from **Image 10.1**, explain why the allele is **not** on the X chromosome. [2]

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- (c) Using **D** for the dominant allele and **d** for the recessive allele, give the genotypes of individuals **5**, **6** and **10**. [2]

5

6

10

- (d) Calculate the probability that individual **9** is a carrier of the allele.

[2]

Probability =

- (e) Explain why at least one of individuals **1** and **2** must carry the allele.

[1]

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11. **Image 11.1** is a light micrograph of a stained blood smear from a person with sickle cell anaemia.

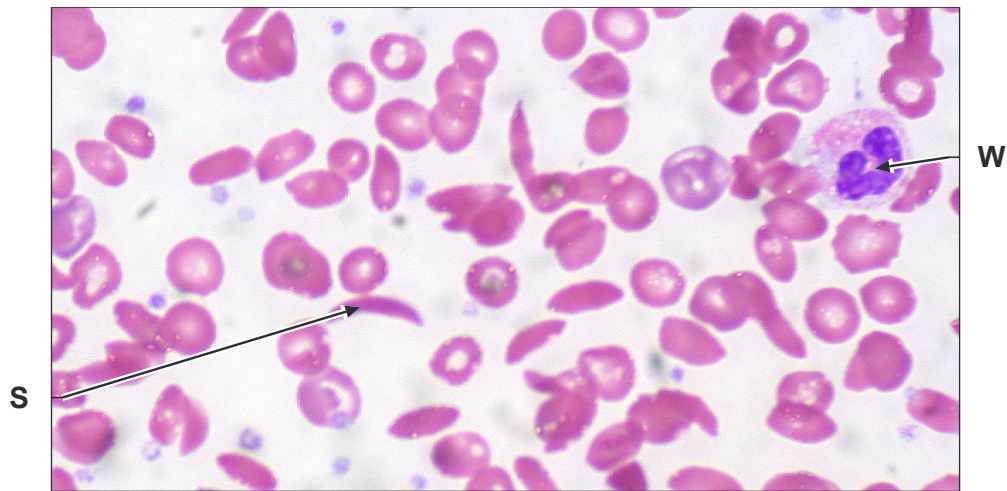


Image 11.1

(a) Identify the cells labelled **S** and **W**. [2]

S

W

Sickle cell anaemia is caused by a change in the gene for the β polypeptide of haemoglobin. **Table 11.2** shows the sixth triplet of the coding strand of DNA for the normal allele, Hb^A , and the sickle cell allele, Hb^S . The mRNA codon GAG codes for glutamic acid and GUG codes for valine.

(b) (i) Complete **Table 11.2**. [2]

Table 11.2

	Hb^A	Hb^S
DNA coding strand	GAG	GTG
mRNA codon		
Amino acid		

(ii) Name the type of gene mutation that produced the Hb^S allele. [1]

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(c) Explain how this change produces cells like **S**. [3]

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- (d) The alleles Hb^A and Hb^S are codominant. Suggest why a heterozygous person usually has only mild symptoms, if any. [2]

12. Duchenne muscular dystrophy (DMD) is caused by recessive alleles of the dystrophin gene on the X chromosome. The gene has 79 exons. Many cases are caused by the deletion of one or more exons. Boys with some deletions have Becker muscular dystrophy instead, a milder condition that starts later in life. **Table 12.1** shows the deletions in two boys.

Table 12.1

Boy	Exons deleted	Number of bases deleted	Condition
P	45	176	DMD
Q	45, 46 and 47	474	Becker muscular dystrophy

- (a) Explain why the deletion in boy **Q** does not change the amino acid sequence after the deleted region. [2]

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- (b) Explain why the deletion in boy **P** results in no functional dystrophin. [3]

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- (c) Suggest why boy **Q** has milder symptoms than boy **P**. [2]

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- (d) Some substitution mutations in the dystrophin gene have no effect on the protein. Explain why. [1]

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(e) State why DMD is extremely rare in girls.

[1]

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13. **Image 13.1** shows meiosis in a cell during the formation of eggs. Only one pair of chromosomes, chromosome 21, is shown.

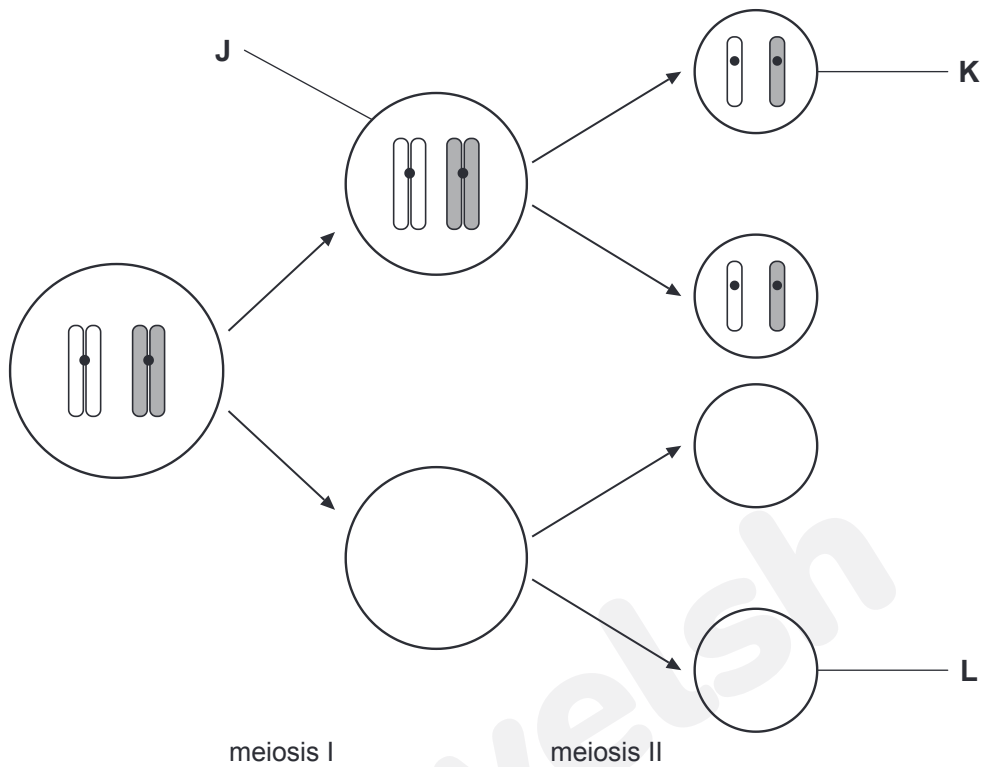


Image 13.1

- (a) Name the error shown in cell **J**, and state the stage of meiosis at which it happened. [2]

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- (b) State the total number of chromosomes in human gametes **K** and **L**. [2]

K

L

- (c) Explain how gamete **K** could lead to a child with Down's syndrome. [2]

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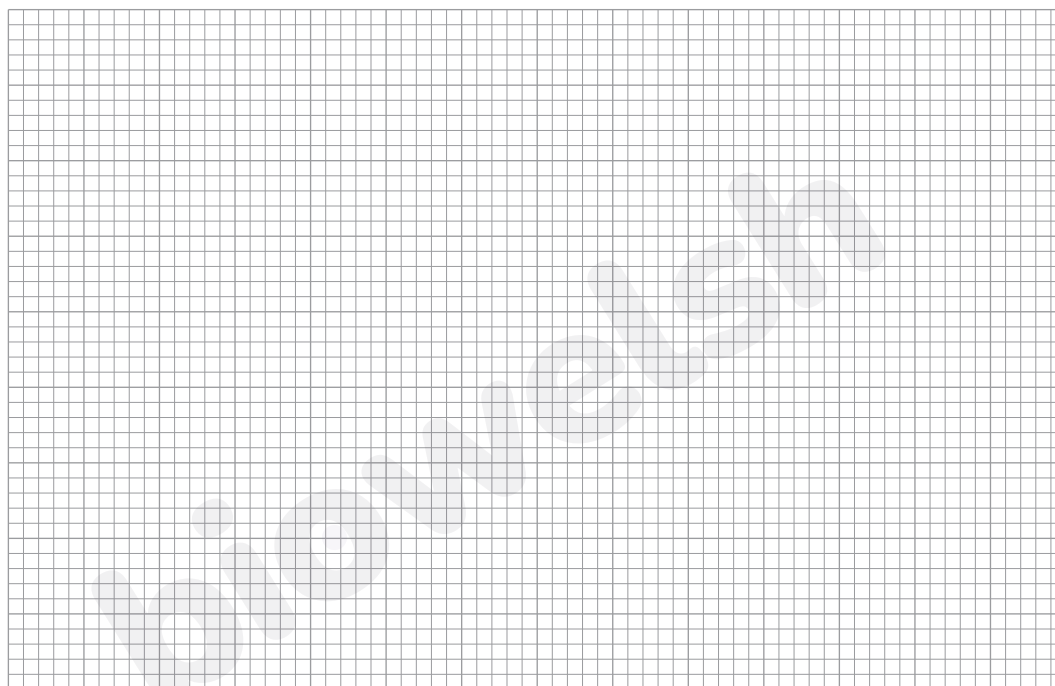
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Table 13.2 shows the number of babies born with Down's syndrome per 1000 births, for mothers of different ages.

Table 13.2

Age of mother / years	20	25	30	35	40	45
Babies with Down's syndrome per 1000 births	0.7	0.7	1.1	2.9	10.0	33.3

- (d) Plot a graph of the data in **Table 13.2** on the grid below. Join the points with straight lines. [4]

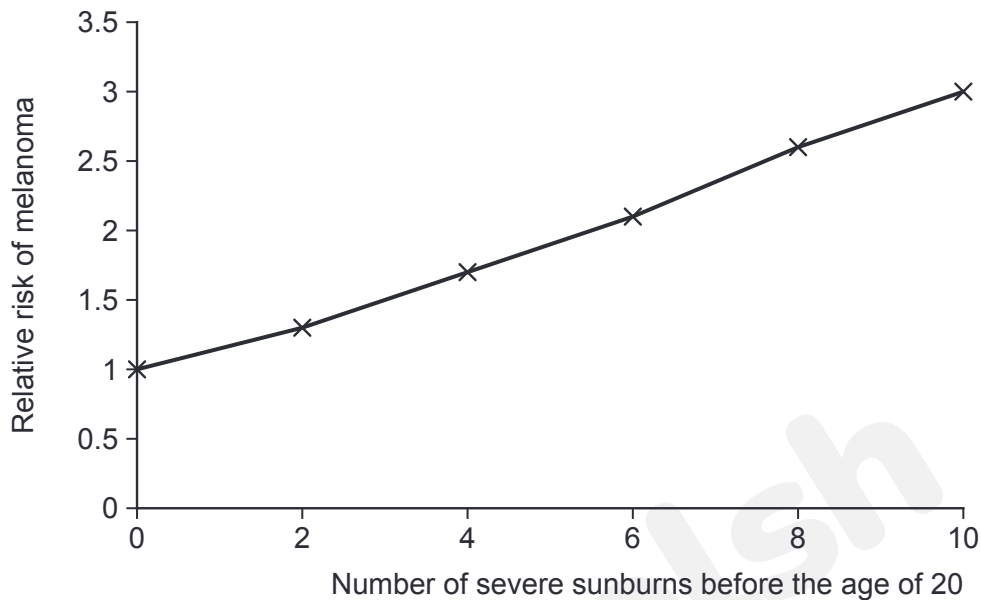


- (e) Suggest why the chance of non-disjunction increases with the age of the mother. [1]

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14. Melanoma is a cancer of the pigment cells in the skin. Researchers recorded the number of severe sunburns that people had before the age of 20, and their later risk of melanoma. **Graph 14.1** shows the results of the study.



Graph 14.1

- (a) Describe the relationship shown in **Graph 14.1**. [2]

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- (b) Calculate the percentage increase in the risk of melanoma for a person with 6 severe sunburns compared with a person with none. [2]

Percentage increase = %

- (c) Ultraviolet light is a carcinogen. Explain how ultraviolet light can cause a melanoma. [3]

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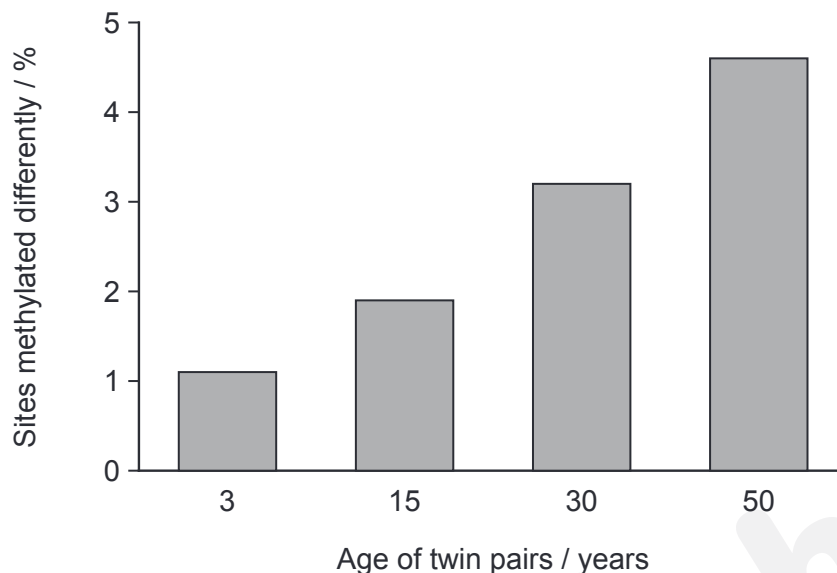
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(d) Explain why these results alone do not prove that sunburn causes melanoma. [2]

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- 15.** Identical (monozygotic) twins develop from the same zygote, so they have the same DNA base sequences. Researchers compared the methylation of DNA at the same sites in the two members of pairs of identical twins of different ages. **Graph 15.1** shows the percentage of sites that were methylated differently in the two twins.



Graph 15.1

Explain what is meant by epigenetics, and describe how DNA methylation and the modification of histones control gene expression. Use **Graph 15.1** to describe the results and suggest explanations for them. Explain how epigenetic changes could lead to one twin developing a cancer that the other twin does not develop. [9 QER]

[9 QER]

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Handwriting practice area with 20 horizontal dotted lines.

END OF WORKSHEET

9





A Tutor Meets Tutee project

BioWelsh worksheets are written and checked by Tutor Meets Tutee tutors for WJEC A level Biology students.

Mark this worksheet with the BioWelsh mark scheme **BW-A2-4.3-W1** (MS).

Acknowledgements

Image 8.1: Photo: National Human Genome Research Institute, public domain (Wikimedia Commons, 'NHGRI human male karyotype'). Letters and label lines added.

Image 11.1: Micrograph: Ed Uthman, CC BY 2.0 (Wikimedia Commons, 'Sickle Cell Anemia (5610746554)'), <https://creativecommons.org/licenses/by/2.0>. Cropped; letters and label lines added.

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