Q1		to blight is a disease of potato plants.	
		to blight is caused by the fungus <i>Phytophthora infestans</i> .	
	(a)	What is the genus of the fungus that causes potato blight?	
	` ,	Tick (✓) one box.	
		Infestans	
		Phytophthora	
		Phytophthora infestans	(1)
	(b)	The fungus grows near the surface of the potato.	(1)
		How does growing near the surface help the fungus to respire?	
		Tick (✓) one box.	
		The fungus can get nitrogen from the soil.	
		The fungus can get oxygen from the air.	
		The fungus can get water from the potato.	
			(1)
	A far	mer sprays his potato plants with a pesticide.	
	The	pesticide kills the fungus that causes potato blight.	
		ying the crop with a pesticide could decrease biodiversity in a river flowing ugh his farm.	
	(c)	What does 'biodiversity in a river' mean?	
		Tick (✓) one box.	
		The variety of species of animals in the river.	

	The variety of species of or the river.	ganisn	ns in			
	The variety of species of plantiver.	ants in	the			(1)
(d)	The farmer sprayed pesticid heavily.	e on h	is potato	plants.	Γhe next day it rained	
	Explain why the biodiversity	in the	river dec	creased.		
Anot	her method of preventing pota	ato bliį	ght is to t	oreed po	tatoes that are resistant	(2)
Resi	stance to potato blight is cont	rolled	by two al	leles:		
	a dominant allele for having re recessive allele for not havir					
A sc	ientist crosses two potato pla	nts. Ea	ach plant	has the	genotype Rr .	
(e)	Complete the diagram below offspring produced.	v to sh	ow the p	ossible g	enotypes of the	
			Male g	ametes		
			R	r		
	Female	R	RR			
	gametes	r				
					_	(2)

(f) Draw a ring around **one** of the homozygous genotypes in the diagram above.

(1)

(g) What percentage of the offspring in the diagram will be resistant to potato blight?

Q2.

	Tick (✓) one box.	
	25% 50% 75% 100%	(1)
(h)	Potatoes can also reproduce asexually.	(-)
	Potatoes from one plant can be planted in the ground to produce new potato plants.	
	All the new plants from a parent plant that is resistant to blight will also be resistant to blight.	
	Explain why.	
		(2)
	(Total 11 n	narks)
The	nucleus of a cell contains DNA.	
(a)	Name the structures inside the cell nucleus that contain DNA.	
		(1)

Figure 1 shows part of a DNA molecule.

T A C G C Phosphate

(b)	Name the part of the DNA molecule labelled X.
(c)	What type of substances are labelled A , C , G and T in Figure 1 ?
Figu	ure 2 shows another section of a DNA molecule.
	Figure 2
	A C T G G C T
(d)	Four of the substances you named in part (c) are not labelled in part Y of Figure 2 .
	Label each of these substances with the correct letter, A, C, G or T.
	Use information from other parts of Figure 2 to help you.
(e)	What is happening to the DNA in part Z of Figure 2 ?
	Tick (✓) one box.
	Differentiation
	Evolution

(Total 8 marks)

AQA Biology GCSE - Reproduction

	Fertilisation		
	Replication		
(f)	A gene is a length of DNA.		(1)
	What type of substance does a gene	code for?	
(g)	Most human body cells contain 6 × 10) ^{–12} grams of DNA.	(1)
	What mass of DNA will a human sper	m cell contain?	
	Tick (✓) one box.		
	6 x 10 ⁻⁶ grams		
	6 × 10 ⁻¹² grams		
	3 × 10 ⁻⁶ grams		
	3 x 10 ⁻¹² grams		
(h)	What is the name of the type of cell di	vision that produces sparm cells?	(1)
(11)	Tick (✓) one box.	vision that produces sperm cens:	
	Binary fission		
	Differentiation		
	Meiosis		
	Mitosis		
			(1)

Q3.

Sickle cell anaemia is an inherited condition that affects red blood cells.

Sickle cell anaemia is caused by a mutation in the gene for haemoglobin. Haemoglobin is the red pigment found in red blood cells.

A person who is homozygous for the normal haemoglobin allele (**H**^A) produces normal red blood cells.

A person who is homozygous for the mutated allele (H^s):

- produces red blood cells with abnormal haemoglobin
- has red blood cells that can form an altered shape
- has sickle cell anaemia and becomes ill.

A person who is heterozygous:

- has both normal and abnormal haemoglobin in the red blood cells
- has sickle cell trait
- is generally healthy but can become ill in certain circumstances.

(a)	Give the reason why a mutation in the gene coding for haemoglobin could be harmful.	
		(1)
(b)	Figure 1 shows some red blood cells from the blood of a person with sickle	

(b) Figure 1 shows some red blood cells from the blood of a person with sickle cell trait.

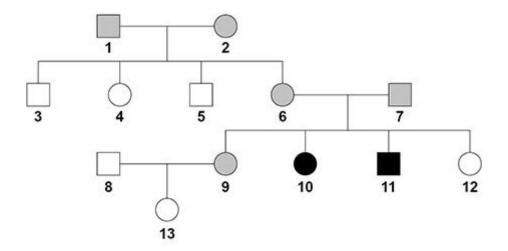
Figure 1



Calculate the proportion of cells in Figure 1 that have an altered sh	ape.
Proportion =	
	(

Figure 2 shows the inheritance of sickle cell anaemia in one family.

Figure 2



Key

- Unaffected male
- Unaffected female
- Male with sickle cell anaemia
- Female with sickle cell anaemia
- Male with sickle cell trait
- Female with sickle cell trait
- (c) Persons 8 and 9 in Figure 2 are expecting a second child.

Determine the probability that the child will be a girl with sickle cell trait.

You should:

- draw a Punnett square diagram
- identify the phenotype of each offspring genotype
- use the symbols:

H^A = normal haemoglobin allele

H^s = mutated haemoglobin allele.

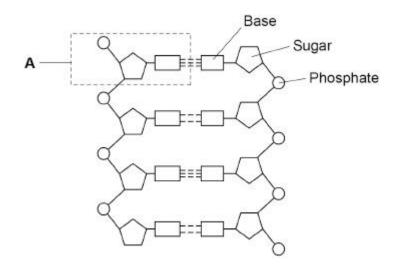
Probability of a girl with sickle cell trait = _____

Q4.

(d)	Without medical treatment, people with sickle cell anaemia are frequently ill and have a reduced life expectancy.
	The malarial parasite cannot live in the red blood cells of a person who has the $\mathbf{H}^{\mathbf{s}}$ allele.
	A scientist stated:
	'It is an advantage for people to have the $\mathbf{H}^{\mathbf{s}}$ allele in countries where malaria occurs.'
	Evaluate the scientist's statement.
	(3) (Total 11 marks)
. Spe	rm cells and egg cells are formed by meiosis.
(a)	During meiosis a cell divides twice.
	How many sperm cells are formed when a cell divides by meiosis?
(b)	(1)
(b)	Human body cells contain 46 chromosomes.
	How many chromosomes are in each human egg cell?
_	(1)
-	uytren's is a disorder that affects the hands.
One	form of Dupuytren's is caused by a dominant allele (D).

The	allele for not ha	ving Dupuytren's	s is recessive	(d).	
(c)	What is an alle	le?			
	Tick (✓) one b	ox.			
	A different for	m of a chromoso	ome		
	A different for	m of a gamete			
	A different for	m of a gene			
(d)	A man with Du	puytren's has th	e genotype D	d.	(1)
	Which word de	scribes the man	i's genotype?		
	Tick (✓) one b	OX.			
	Heterozygous				
	Homozygous				
	Phenotype				
	man with Dupuy plan to have a c		a woman who	o does not have	(1) Dupuytren's
(e)	Complete the g		in the figure b	pelow to show the	possible
			W	oman	_
			d	d	
	Man	D	Dd		
		d	0		
					(2)
(f)	Draw a ring ard have Dupuytre		pe of a child in	n the figure abov	e who will
					(1)

(g)	What is the chance of the child having Dupuytren's?	
	Tick (✓) one box.	
	25%	
	50%	
	75%	
	100%	
		(1)
(h)	A genetic disorder develops as a result of a change in a gene.	
	What scientific term describes a change in a gene?	
(i)	People with a family history of some genetic disorders are offered embryo screening. Suggest one way embryo screening can help people with a family history of a genetic disorder.	
	(Total 10 r	(1) narks)
Q5. Thi	is question is about DNA.	
(a)		
. ,		
		(2)
The	e below diagram shows part of a DNA molecule.	



(b) DNA codes for a sequence of amino acids.

Which part of DNA forms the code for a particular amino acid?

	Tick (✓) one bo	X.
	Bases	
	Phosphates	
	Sugars	
(c)	Which substance	e is produced when amino acids are joined together?
	Tick (✓) one bo	X.
	Carbohydrate	
	Fat	
	Protein	

(1)

(1)

(d) DNA is made of repeating units. One of the units is labelled **A** in the diagram above.

What is the name of the repeating unit labelled **A**?

Tick (\checkmark) one box.

	Length = million nm
Give your ans	Length = million nm swer to question (e) in metres.
•	
•	
•	swer to question (e) in metres.
•	swer to question (e) in metres.
1 metre = 1 >	swer to question (e) in metres. × 10 ⁹ nanometres Length = m
1 metre = 1 >	swer to question (e) in metres. × 10 ⁹ nanometres
1 metre = 1 >	swer to question (e) in metres. × 10 ⁹ nanometres Length = m
1 metre = 1 > DNA analysis Patients who	swer to question (e) in metres. × 10 ⁹ nanometres Length = m s can show people which alleles they have.

(1)

(4)

(b)

	^
()	h
w	u.

DNA is a polymer of nucleotides.

(a) Why is DNA described as a polymer?

The diagram below shows part of a DNA molecule.

Describe the structure of a nucleotide.

(c) The length of a DNA double helix increases by 0.34 nm for every pair of nucleotides.

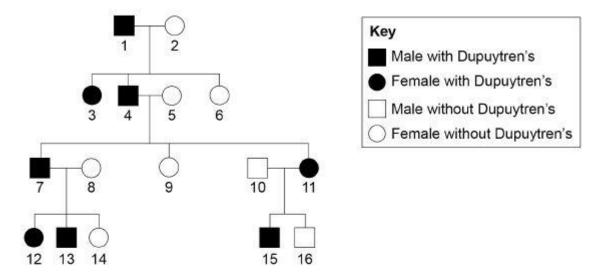
Q7.

Give your answer in metres. Use information from the diagram.
Total length = m
<u></u>
Some parts of DNA do not code for proteins.
Describe how non-coding parts of DNA can affect the expression of genes.
geneer and one of the second of geneer
(Total 11 r
e are two types of cell division: mitosis and meiosis.
Describe three differences between the processes of mitosis and meiosis.
1

Describe on	e similarity be	etween the	processes	of mitosis an	d meiosis.

The diagram halow shows the inheritance of Dupuntron's in one for

The diagram below shows the inheritance of Dupuytren's in one family.



Dupuytren's is caused by a dominant allele in this family.

D = dominant allele

d = recessive allele

(c) Give the genotype of person 1.

Explain your answer.

Genotype _____

(2)

(d) Person **7** and person **8** in the diagram above are expecting a fourth child.

What is the probability of the child having Dupuytren's?

You should:

- draw a Punnett square diagram
- identify which offspring have Dupuytren's

	Probability =	_ (5)
(e)	Explain how the diagram above shows the allele for Dupuytren's is not on the Y chromosome.	
		(2)
	(Total 13	

Q8.

The shape of a person's earlobes is controlled by a gene.

Figure 1 shows two types of earlobe.

Free earlobe

Figure 1

Attached earlobe

A dominant allele codes for free earlobes.

(a) What is a dominant allele?

Tick (✓) one box.

An allele expressed even if a person only has one copy of the allele

An allele expressed only if a person has two copies of the allele

An allele expressed only if a person has no recessive allele

An allele expressed only if it is inherited from the male parent

(1)

(b) A man with free earlobes and a woman with attached earlobes have children together.

Complete Figure 2 to show the possible genotypes of the children.

Use the symbols:

E = allele for free earlobes

e = allele for attached earlobes

Figure 2

Woman
e e

E Ee

Man
e

(2)

(c) What is the probability that one of the children would have attached earlobes?

Use Figure 2.

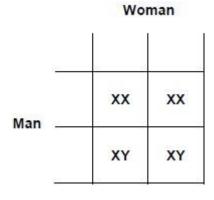
Tick (✓) one box.

0.125	0.25	0.5	0.75	
				(1)

(d) **Figure 3** shows the inheritance of the sex chromosomes, **X** and **Y**.

Complete **Figure 3** to show the sex chromosomes in the gametes of the man and the woman.

Figure 3



(2)

(e) Calculate the probability that the man and the woman's next child will be a girl with attached earlobes.

Use the equation:

probability of a girl with attached earlobes

= probability of attached earlobes × probability of being a girl

Probability of a girl with attached earlobes =_____

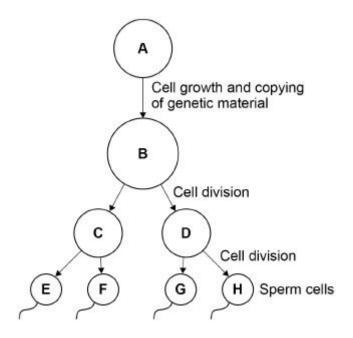
(2)

(Total 8 marks)

Q9.

Figure 1 shows the production of sperm cells in humans.

Figure 1



(a) Cell A is a normal body cell.How many chromosomes are there in cell A?Tick (√) one box.

23	46	0 0	48	0 0	92	0 0	

(b) What is the mass of DNA in cell E?

Tick (\checkmark) one box.

A quarter of the mass of the DNA in cell A	6 /
Half the mass of the DNA in cell A	
The same mass as the DNA in cell A	
Twice the mass of the DNA in cell A	

(1)

(1)

(c) What type of cell division produces sperm cells?

Tick (\checkmark) one box.

	Binary fission		
	Differentiation		
	Meiosis		(1)
(d)	Sometimes there a	are errors in copying the genetic material.	
	What term describ	es an error in the genetic material?	
	Tick (✓) one box.		
	Absorption		
	Fertilisation		
	Mitosis		
	Mutation		(4)
(e)	A woman has thre	e children, aged 4, 6 and 9 years.	(1)
	Why are the childr	en not genetically identical?	
In se cell.	exual reproduction, a	a sperm cell fuses with an egg cell to form a new single	(2)
An e	mbryo develops fro	m the single cell.	
The	cell divides three tin	nes to produce the embryo.	
(f)	How many cells are	e there in the embryo after three cell divisions?	
	Tick (✓) one box.		

(3)

	3	6	8	9	
					(1)
Figure 2	2 shows a differe	ent human embr	yo.		
		Figi	ure 2		
				Ť	
	1	(Decree			

(g) Measure image length X on Figure 2.

Give your answer in millimetres (mm).

(h) The image in **Figure 2** has been magnified × 500

Calculate the real length of the embryo.

Use the equation:

$$real length of the embryo = \frac{image length}{magnification}$$

Give your answer in micrometres (µm).

Real length of the embryo = μ m

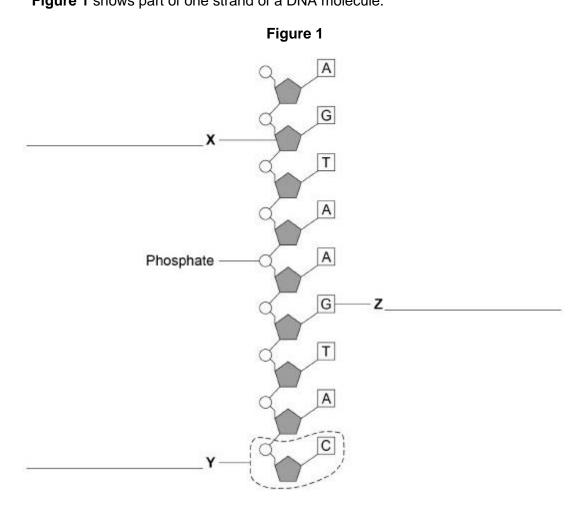
(1)

(i)	The embryo may not implant in the lining of the uterus.
	The embryo will then be lost from the woman's body several days later.
	Explain why the woman may not notice this has happened.
	(2) (Total 13 marks)
Q10.	
∝ 10.	

Figure 1 shows part of one strand of a DNA molecule.

Which structures in the nucleus of a human cell contain DNA?

Genetic material is made of DNA.



(Total 8 marks)

a complete DNA molecule is made of two strands twisted around each other. What scientific term describes this structure?	h
What scientific term describes this structure?	
DNA codes for the production of proteins.	
A protein molecule is a long chain of amino acids.	
How many amino acids could be coded for by the piece of DNA show Figure 1?	n in
Tick (√) one box.	
2 3 9 18	
Scientists have now studied the whole human genome.	
Give two benefits of understanding the human genome.	
1	
2	

Q11.The following table gives the classification of four plant species.

Group	Species 1	Species 2	Species 3	Species 4
Kingdom	Plantae	Plantae	Plantae	Plantae
Phylum	Spermatophyta	Spermatophyta	Spermatophyta	Spermatophyta
Class	Monocotyledonae	Dicotyledonae	Monocotyledonae	Dicotyledonae
Order	Poales	Fabales	Poales	Scrophulariales
Family	Cyperaceae	Fabaceae	Poaceae	Scrophulariaceae

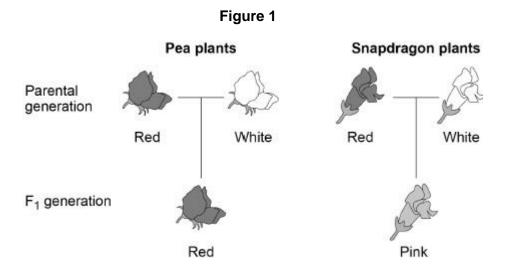
(1)

Genus	Eriophorum	Pisum	Poa	Antirrhinum
Species	angustifolium	sativum	annua	majus

(a) Species 1 and 3 are the most closely related.

Figure 1 shows the inheritance of flower colour in two species of plant.

What information in the table above gives evidence for this?



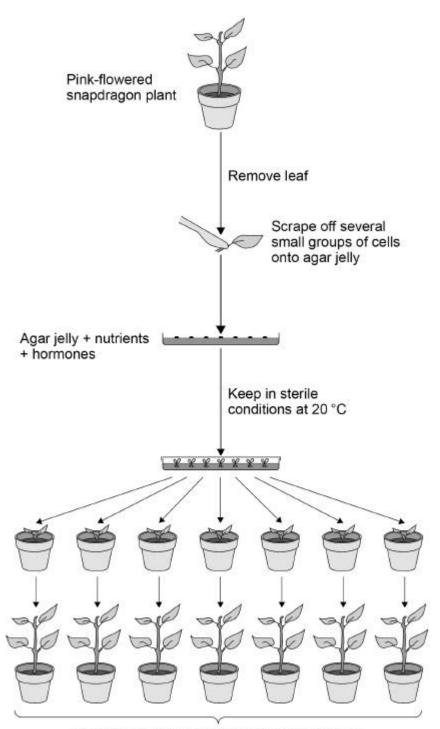
- In pea plants and in snapdragon plants, flower colour is controlled by one pair of alleles.
- In **Figure 1** the parental generation plants are homozygous for flower colour.
- In heterozygous **pea** plants, the allele for red flower colour is dominant.
- In heterozygous snapdragon plants, the alleles for flower colour are both expressed.

Use the following symbols for alleles in your answers to parts (b) to (d):

Pea	plants	Snapdragon plants	
	allele for red flowers llele for white flowers	C^R = allele for red flowersC^W = allele for white flowers	
(b)	What is the genotype of the red-flo	wered pea plants in the F ₁ generation?	
			(1)

(c)	What is the genotype of a white-flowered snapdragon plant?	
		(1)
A ga	ardener crossed two pink-flowered snapdragon plants.	
(d)	Draw a Punnett square diagram to show why only some of the next generation plants had pink flowers.	
	Identify the phenotypes of all the offspring plants.	
		(3)
(e)	What percentage of the offspring would you expect to have pink flowers?	
	Percentage =%	(1)
	nmercially, hundreds of pink-flowered snapdragon plants can be produced none pink-flowered plant.	(1)
	ure 2 shows a tissue culture technique used for producing many plants from plant.	

Figure 2



Many snapdragon plants, all with pink flowers

(f) Give a reason for each of the following steps shown in Figure 2.Several groups of cells are scraped off the leaf:

Nutrients are added to the agar jelly: _____

	Hormones are added to the agar jelly:	
	The plant cells are kept in sterile conditions:	
	The plant cells are kept at 20 °C:	-
		(
(g)	Explain why the method shown in Figure 2 produces only pink-flowered plants.	
		,
	(Total 14	(mark
12. This	question is about the cell cycle.	
(a)	Chromosomes are copied during the cell cycle.	
	Where are chromosomes found?	
	Tick one box.	
	Cytoplasm	
	Nucleus	

	Ribosomes		
	Vacuole		
			(1)
(b)	What is the nam characteristic?	e of a section of a chromosome that controls a	
			(1)

Figure 1 shows information about the cell cycle.

Mitosis

Cell growth

Copying of chromosomes

Figure 1

(c) Which stage of the cell cycle in **Figure 1** takes the most time?

Tick **one** box.

Cell growth

Copying of chromosomes

Mitosis

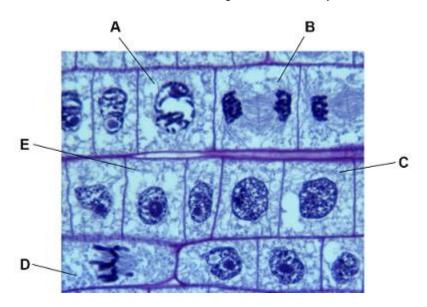
(1)

(d) During mitosis cells need extra energy.Which cell structures provide most of this energy?

Tick one box.

		Time = minutes	(2
	Calculate the tim	ne taken for mitosis.	
(e)		Figure 1 takes two hours in total. stage takes 45 minutes.	
			(1
	Ribosomes		
	Mitochondria		
	Cytoplasm		
	Chromosomes		

Figure 2 shows some cells in different stages of the cell cycle.



(f) Which cell is **not** dividing by mitosis

Tick one box.

	A B C D	(1)
(g)	Cell E in Figure 2 contains 8 chromosomes. Cell E divides by mitosis.	()
	How many chromosomes will each new cell contain?	
	Tick one box.	
	2	
	4	
	8	
	16	
		(1)
(h)	Why is mitosis important in living organisms?	
	Tick one box.	
	To produce gametes	
	To produce variation	
	To release energy	
	To repair tissues	
		(1) (Total 9 marks)

Q13.

In the mid-19th century, a scientist studied inheritance in pea plants.

The scientist's work was the beginning of our modern understanding of genetics.

(a) What is the name of this scientist?

Tick **one** box.

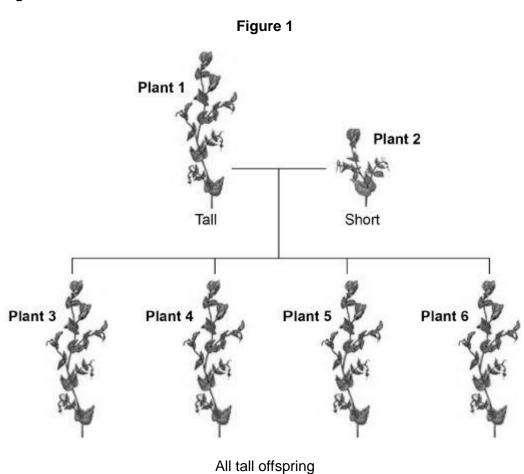
	Alfred Russel Wallace	е	
	Charles Darwin		
	Gregor Mendel		
	Jean-Baptiste Lamar	ck	
(b)	In the mid-20th centur that makes up genetic	y, other scientists identified the chemical substance material.	(1)
	What is the name of the material?	ne chemical substance that makes up genetic	
	Tick one box.		
	Carbohydrate		
	DNA		
	Lipid		
	Protein		
			(1)
(c)	A gene often has two	alleles.	
	One allele is dominan	t and the other allele is recessive.	
	When is a recessive a	llele expressed as a characteristic?	
	Tick one box.		
	When the dominant a	Illele is not present	
	When the recessive a	allele is inherited from the female parent	
	When the recessive a	allele is inherited from the male parent	

When the recessive allele is present on only one of the chromosomes (1)

A scientist investigated the inheritance of height in pea plants.

The scientist crossed tall pea plants with short pea plants.

Figure 1 shows the scientist's results.



In questions (d) and (e), use the following symbols to represent alleles:

T = the dominant allele for tall.

t = the recessive allele for short.

(d) In Figure 1, the genotype of plant 1 is TT.

Give the genotype of plant 2.

(1)

(e) The scientist crossed plant 3 with plant 4.

Complete Figure 2 to show the offspring produced from this cross.

Figure 2

		Male gametes	
		Т	t
Female	T ***	тт	٠
gametes	t		

(2)

(f) Draw a circle around **one** of the homozygous offspring in **Figure 2**.

(1)

(g) What is the ratio of tall plants: short plants in the offspring in Figure 2?

Datio of tall plants , short plants	•
Ratio of tall plants : short plants =	_ •

(1)

(Total 8 marks)

Q14.

Cell division is needed for growth and for reproduction.

(a) The table below contains three statements about cell division.

Complete the table.

Tick **one** box for each statement.

	Statement is true for		e for
Statement	Mitosis only	Meiosis only	Both mitosis and meiosis
All cells produced are genetically identical			
In humans, at the end of cell division each cell contains 23 chromosomes			
Involves DNA replication			

(2)

Bluebell plants grow in woodlands in the UK.

- Bluebells can reproduce sexually by producing seeds.
- Bluebells can also reproduce asexually by making new bulbs.

þ	arent is needed.	
S	Suggest two other advantages of asexual reproduction for bluebells.	
1	·	
2	·	
_		
Ε	xplain why sexual reproduction is an advantage for bluebells.	
_		
_		
_		
_		
_		
_		
_	······································	
-	_	

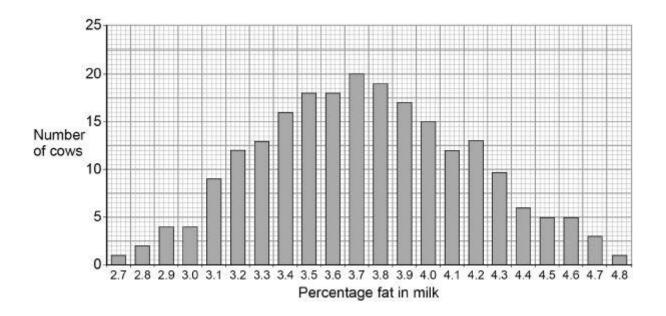
Q15.

Scientists want to breed cows that produce milk with a low concentration of fat.

Figure 1 shows information about the milk in one group of cows.

The cows were all the same type.

Figure 1



(a) In **Figure 1** the mean percentage of fat in the milk is equal to the modal value.

Give the mean percentage of fat in the milk of these cows.

Mean percentage = ______(1)

(b) A student suggested:

'The percentage of fat in milk is controlled by one dominant allele and one recessive allele.'

How many different phenotypes would this produce?

Tick one box.

2 3 22 46

(1)

(c) Give the evidence from **Figure 1** which shows the percentage of fat in the milk is controlled by several genes.

(1)

(d) One of the genes codes for an enzyme used in fat metabolism.

A mutation in this gene causes a reduction in milk fat.

The mutation changes one amino acid in the enzyme molecule.

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enzyme working.		
	 	

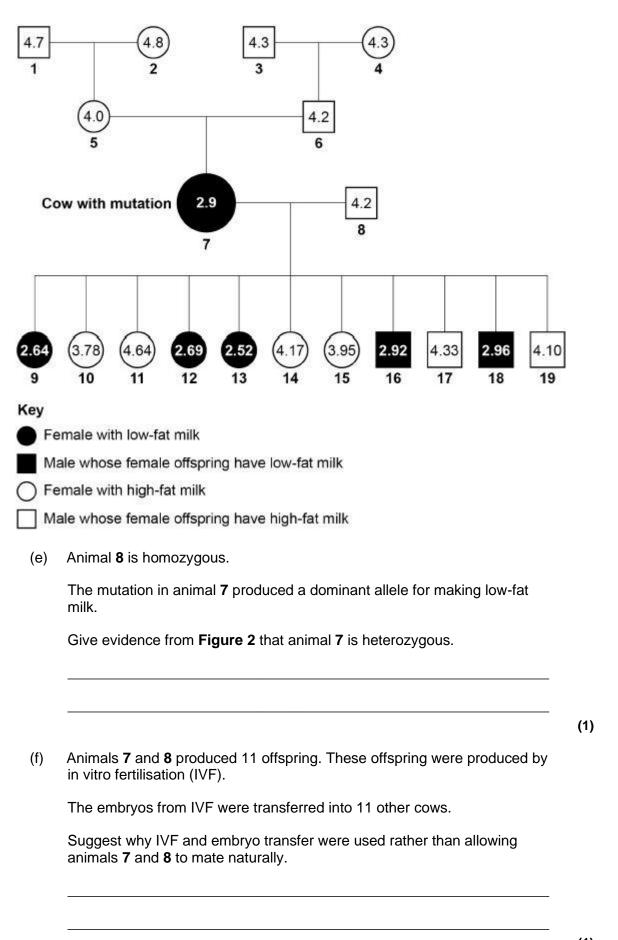
The scientists found one cow with a mutation.

The cow's milk contained only 2.9% fat.

Figure 2 shows the percentage of fat in the milk of cattle related to the cow with the mutation.

The values for male cattle are the mean values of their female offspring.

Figure 2



Draw a Punnett square diagram to show a cross between animals 7 and 8.	
Identify which offspring produce low-fat milk and which offspring produce high-fat milk.	
Use the following symbols:	
D = dominant allele for making low-fat milk	
d = recessive allele for making high-fat milk	
	(4)
	(+)
The scientists want to produce a type of cattle that makes large volumes of low-fat milk.	
The scientists will selectively breed some of the animals shown in Figure 2 .	
Describe how the eciontists would do this	
Describe now the scientists would do this.	
	(4)
(Total 16 m	
	Identify which offspring produce low-fat milk and which offspring produce high-fat milk. Use the following symbols: D = dominant allele for making low-fat milk d = recessive allele for making high-fat milk The scientists want to produce a type of cattle that makes large volumes of low-fat milk.

$\overline{}$	4	^
u	1	b.

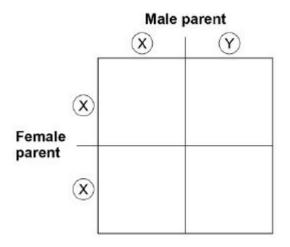
m	nitosis ov	ary	sperm	testis	uterus
exu	ual reproduction	binary fiss	ion egg	fertilisation	meiosis
	Choose the answ	ers from the b	OOX.		
c)	Complete the ser	tences.			
b)	How many chrom	osomes are t	here in a huma	n gamete cell?	
	48				
	46				
	24				
	23				
	Tick one box.				
a)	How many chrom	osomes are t	here in most hu	ıman body cells?	
	omosomes are four	,			

The female gamete is called the
The male gamete is called the
The female gamete is produced in the
Gametes are produced by a type of cell division
called
Male and female gametes join together in a process
called

(5)

In humans, the sex chromosomes are called **X** and **Y**.

The diagram shows the inheritance of sex chromosomes.



(d)	Complete the diagram above to show the sex chromosomes inherited by
	the offspring.

(2)

(e) What is the chance that a child produced by these parents will be female?Tick one box.

1 in 2

1 in 3

1 in 4

3 in 4

(1)

(f) The parents shown in the diagram above have five children.

Give **two** reasons why these children all look different from each other.

1. ______

2. _____

(2)

		(Total 12 ma	rks)
Q17		mans, chromosome X and chromosome Y are the sex chromosomes.	
	a)	Most cells in the human body contain two sex chromosomes.	
		Which type of cell does not have two sex chromosomes?	
		Tick one box.	
		Liver cell	
		Muscle cell	
		Nerve cell	
		Red blood cell	
(b)	Apart from the sex chromosomes, how many other chromosomes are there in most human body cells?	(1)
		Tick one box.	
		21 23 44 46	(1)
		Stickler syndrome is an inherited disorder that causes damage to the eye.	(-)
		One of the symptoms of Stickler syndrome is that black spaces can appear in the visual image.	
(c)	Which part of the eye is affected by Stickler syndrome?	
		Tick one box.	
		Ciliary muscles	
		Iris	

Retina

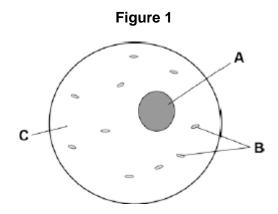
(Total 9 marks)

	Suspensory ligaments	(1)
Stick	kler syndrome is caused by the inheritance of a dominant allele.	(-)
The	diagram shows the inheritance of Stickler syndrome in two families.	
13	1 2 3 4 Key: = affected m = affected fee = unaffected = unaffected = unaffected = unaffected	male I male
$\mathbf{A} = \mathbf{t}$	the following symbols in your answers to (d) and (e): the dominant allele for Stickler syndrome the recessive allele for unaffected vision. Explain why none of the children of persons 7 and 8 have Stickler syndrome.	
(e)	Person 12 marries person 18. Use a Punnett square diagram to find the probability that their first child will be a female with Stickler syndrome.	(2)
	Probability of a female child with Stickler syndrome =	(4)

(2)

Q18.

Figure 1 shows a human body cell.



(a) Which part in Figure 1 contains chromosomes?

Tick one box.

(b) Humans have pairs of chromosomes in their body cells.

Draw **one** line from each type of cell to the number of chromosomes it contains.

Type of cell	Number of Chromosomes
	10
Human body cell	23
	46
Sperm cell	60
	92

(c) Humans have two different sex chromosomes, **X** and **Y**.

Figure 2 shows the inheritance of sex in humans.

Figure 2

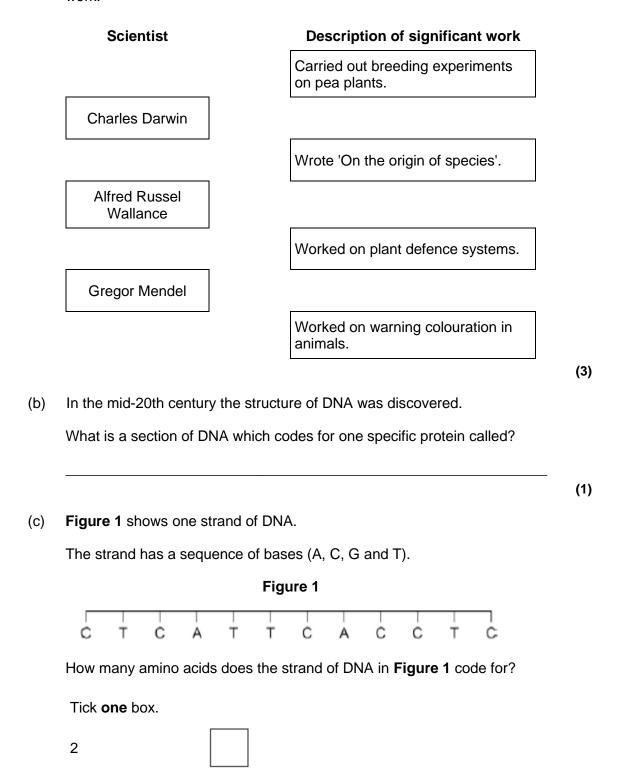
rcle a part of Figure 2 that shows a ve the genotype of male offspring. man and a woman have two sons. ild. hat is the chance that this child will ick one box. 0%	The womar	n is p	regnant	with a third
man and a woman have two sons. ild. hat is the chance that this child will ick one box.	The womar	n is p	regnant	with a third
man and a woman have two sons. ild. hat is the chance that this child will ick one box.			regnant	with a third
ild. hat is the chance that this child will ick one box.			regnant	with a third
ild. hat is the chance that this child will ick one box.			regnant v	with a third
ick one box.	also be a b	ooy?		
0%				
25%				
50%				
00%				

Mother

Q19.

Our understanding of genetics and inheritance has improved due to the work of many scientists.

(a) Draw **one** line from each scientist to the description of their significant work



	3					
	4					
	6					
						(1)
(d)	Mutations of DNA cause some	inhei :	rited d	isorde	ers.	
	One inherited disorder is cystic	c fibro	sis (C	F).		
	A recessive allele causes CF.					
	Complete the genetic diagram	in Fi ç	gure 2			
	Identify any children with	ı CF.				
	Give the probability of an	ny chil	dren h	naving	g CF.	
	Each parent does not have CF.					
	The following symbols have been used:					
	D = dominant allele for not ha	ving C	CF			
	d = recessive allele for having	CF				
		Figur	e 2			
		1	Mothe	r		
			D	d	_	
	Father	D	DD			
		d				
	Probability o	of a ch	ild with	n CF :	=	
						(3)
(e)	What is the genotype of the m	other	showr	ı in Fi	gure 2?	
	Tick one box.					
	Heterozygous					
	Homozygous dominant					

Homozygous recessive		
		(1)
		(Total 9 marks)

Q20.

Figure 1 shows an image of a small section of DNA.

Figure 2 shows the structure of a small section of DNA.

Figure 1 Figure 2 Part B Bases © Svisio/iStock/Thinkstock

What is Part B? (a)

(1)

(b) In **Figure 1** the structure of DNA shows four different bases.

There are four different bases and they always pair up in the same pairs.

Which bases pair up together?

(1)

(c) Syndrome H is an inherited condition.

People with syndrome H do **not** produce the enzyme IDUA.

Figure 3 shows part of the gene coding for the enzyme IDUA.

Figure 3

(Total 12 marks)

	Ċ	Å	Ť	T	С	A	G	C	T	С	Strand J from a person without syndrome H
T	C	A	T	T	T	A	G	С	T	С	Strand K from a persor with syndrome H
	Strand K shows a mutation in the DNA which has caused syndrome H.										sed syndrome H.
	The enzyme IDUA helps to break down a carbohydrate in the human body.										
	The e	enzyn	ne IDU	JA pro	oduce	d fron	n Stra	nd K \	will no	t wo	rk.
	Expla	ain ho	w the	muta	tion c	ould c	ause	the er	nzyme	not	to work.
(d)	A rec	cessiv	e allel	le cau	ses s	yndro	me H.				
(d)		terozy							recess	sive r	man want to have a
(d)	A hetchild.	terozy	gous nnett	woma	an and	d a ho	mozy	gous ı			
(d)	A her child. Draw havir	terozy / a Pu	gous nnett	woma squar e H.	an and e diaç	d a ho gram t	mozy to dete	gous ı			man want to have a
(d)	A her child. Draw havir	terozy / a Pu ng syn	rgous Innett Indromo y child	woma squar e H. dren w	an and e diaç	d a ho gram t	mozy to dete	gous ı			man want to have a
(d)	A her child. Draw havir Ident	terozy · · a Pu ng syn tify an	rgous nnett ndrom y child	woma squar e H. dren w g sym	an and e diaç	d a ho gram t	mozy to dete	gous ı			man want to have a
(d)	A her child. Draw havir Ident Use t	terozy	rgous nnett ndrome y child llowing ant all	woma squar e H. dren w g sym	an and e diaç	d a ho gram t	mozy to dete	gous ı			man want to have a

0	2	1	
w	_		

In humans, hair colour is an inherited characteristic.

Red hair is caused by a recessive allele.

(a) When does a recessive allele control the development of a characteristic?

Tick (**√**) **one** box.

When the allele is present on only one of the chromosomes.

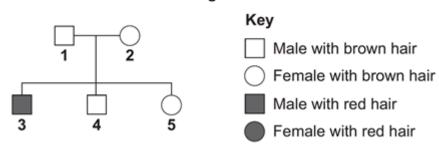
When the dominant allele is not present.

When the allele is inherited from the female parent.

(1)

(b) **Figure 1** shows the inheritance of hair colour in one family.

Figure 1



(i) Brown hair is caused by a dominant allele, **B**.

Red hair is caused by the recessive allele, **b**.

What combination of alleles does person 1 have?

Tick (**√**) **one** box.

ВВ

Bb

bb

(1)

(ii) Person 3 married a woman with brown hair.

Figure 2 shows how hair colour could be inherited by their children.

Figure 2

		Woman Brown hair				
		В	b			
Person 3	b	Bb				
Red hair	b					

Complete **Figure 2** to show the combination of alleles that the children would inherit.

One has been done for you.

(2)

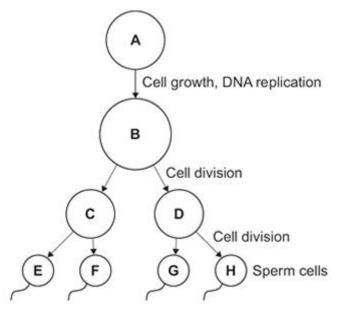
(iii) What is the probability that one of the children would have red hair?
Tick (✓) one box.
1 in 2
1 in 3
1 in 4

(1)

(Total 5 marks)

Q22.

The diagram below shows the production of human sperm cells.



(a)	Name the organ where the processes shown in the diagram above take place.						
(b)	(i)	Not every cell in the diagram above contains the same amount of DNA.	(1)				
		Cell A contains 6.6 picograms of DNA (1 picogram = 10 ⁻¹² grams).					

How much DNA is there in each of the following cells?

Cell **B** ______ picograms

Cell **C** _____ picograms

Cell **E** _____ picograms

(2)

(ii) How much DNA would there be in a fertilised egg cell?

pic	cograms
	(1)

(iii) A fertilised egg cell divides many times to form an embryo.

Name this type of cell division.

(1)

•	What are stem cells?
-	
_	
Į	Suggest why it is ethically more acceptable to take stem cells from a umbilical cord instead of using stem cells from a 4-day-old embryo produced by In Vitro Fertilisation (IVF).
	Stem cells taken from a child's umbilical cord could be used to treat a condition later in that child's life.
	Give one advantage of using the child's own umbilical cord stem cells instead of using stem cells donated from another person.
	Why would it not be possible to treat a genetic disorder in a child using his own umbilical cord stem cells?
-	

Q23.

Polydactyly is an inherited condition caused by a dominant allele.

(a) The figure below shows the hand of a man with polydactyly. The man has an extra finger on each hand.

The man's mother also has polydactyly but his father does not.



© Ifness/iStock

neteroz	how the information given above shows that the man is rygous for polydactyly.
The ma	n marries a woman who does not have polydactyly.

(b)

Brown hair is caused by th	e dominant allele,	В.	
Red hair is caused by a re	cessive allele, b .		
Complete the genetic diag able to have some children			
	Father	Mother	
Parental phenotypes			
Parental Genotypes			
Gametes			
0"			
Offspring genotypes:			
Offspring phenotypes:			
			(5) (Total 9 marks)
			•

The man has red hair. His sister has brown hair.

Both of their parents have brown hair.