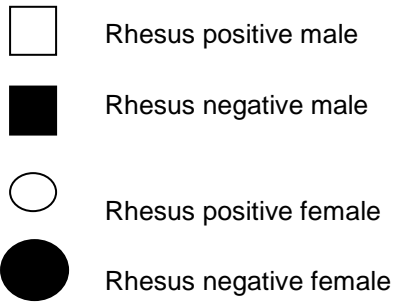
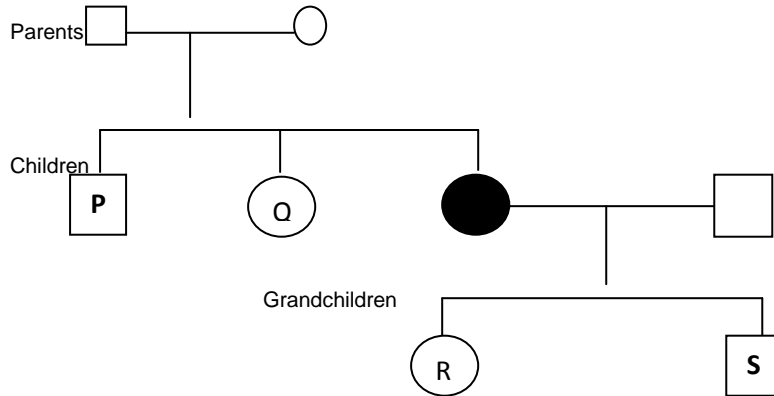


### Reproduction 3

1. The family tree below shows the transmission of the Rhesus D-antigen through three generations of a family. The allele coding for the presence of the Rhesus D-antigen is dominant and autosomal.

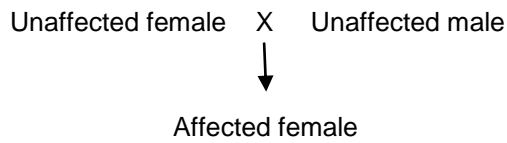


Which of the children and grandchildren in the family tree must be heterozygous?

- A P, Q, R and S
- B P and Q only
- C R and S only
- D Q and R only

Reproduction 3

2. The family tree shows the pattern of inheritance of a genetic condition.



The allele responsible for this condition is both

- A** sex –linked and recessive  
**B** sex – linked and dominant  
**C** autosomal and recessive  
**D** autosomal and dominant
3. Haemophilia is a sex-linked recessive condition. A woman, who does not have this condition, has a haemophiliac son. The boy's father is also a haemophiliac.

What are the genotypes of the parents?

	Father	Mother
<b>A</b>	$X^H Y$	$X^H X^h$
<b>B</b>	$X^h Y$	$X^h X^h$
<b>C</b>	$X^h Y$	$X^H X^H$
<b>D</b>	$X^h Y$	$X^H X^h$

4. Cystic fibrosis is a genetic condition caused by an allele which is not sex-linked.

A child is born with cystic fibrosis despite neither parent having the condition.

The parents are going to have a second child. What is the chance this child will have cystic fibrosis?

- A.** 1 in 2   **B.** 1 in 3   **C.** 1 in 4   **D.** None
5. Huntington's Disease is an inherited condition in humans caused by a dominant allele which is not sex-linked.

A woman's father is heterozygous for the condition and her mother is unaffected.

What is the chance of the woman having the condition?

- A** 1 in 1  
**B** 1 in 2  
**C** 1 in 3

Reproduction 3

D 1 in 4

6. A sex-linked condition in humans is caused by a recessive allele. What is the chance of an unaffected man and a carrier woman having an unaffected male child?

A 1 in 1

B 1 in 2

C 1 in 3

D 1 in 4

7. The table below shows some genotypes and phenotypes associated with a form of anaemia

<i>Genotype</i>	<i>Phenotype</i>
AA	Unaffected
AS	Sickle cell trait
SS	Acute sickle cell anaemia

An unaffected person and someone with sickle cell trait have a child together.

What are the chances of the child having acute sickle cell anaemia?

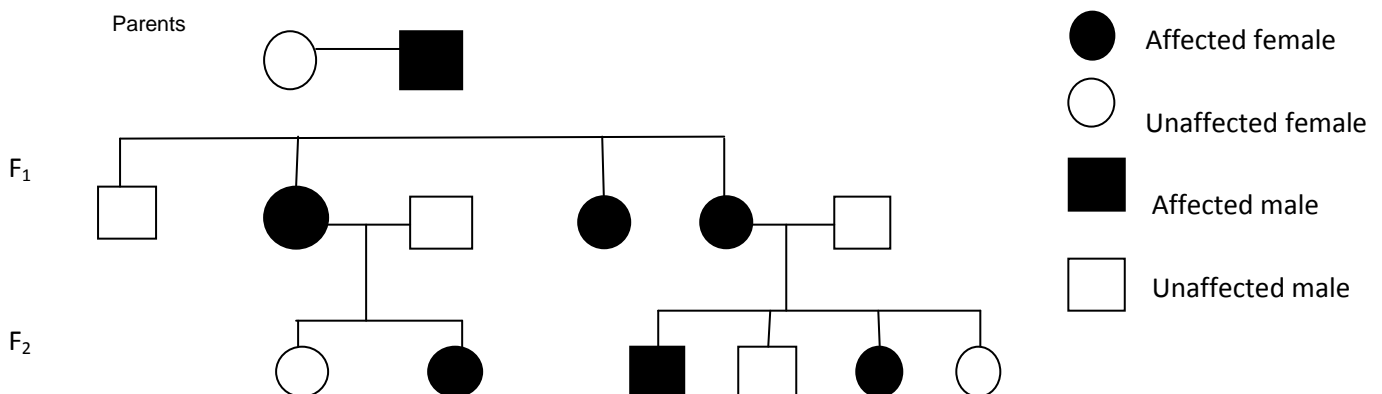
A none

B 1 in 4

C 1 in 2

D 1 in 1

8. The family tree shows the inheritance of a bone disorder.



The disorder is caused by a dominant sex-linked allele (B).

(a) Using appropriate symbols, give the genotypes of individuals P, Q, R, S.

P \_\_\_\_\_ Q \_\_\_\_\_ R \_\_\_\_\_ S \_\_\_\_\_

Reproduction 3

(b) (i) Explain why all the  $F_1$  females in this family are affected.

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(ii) Explain why only some of the  $F_2$  females in this family are affected.

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(c) Is the ratio of affected offspring to unaffected offspring in the  $F_1$  generation as expected? Give a reason for your answer.

Yes/No \_\_\_\_\_

Reason \_\_\_\_\_

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