

#### **Normal Vision**



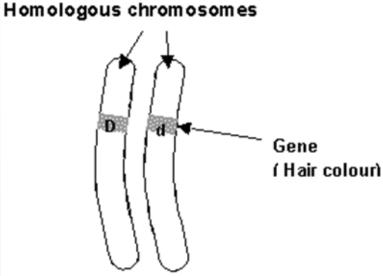
#### Colourlblind

S1-1-14 Explain the inheritance of sex-linked traits in humans and use a pedigree to track the inheritance of a single trait. Examples: colour blindness, hemophilia

#### **Genes & Alleles...**

#### <u>Genes</u>

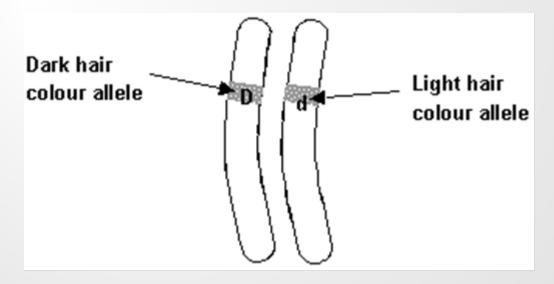
- Genes are a <u>PIECE</u> of <u>CHROMOSOME</u> that contains the actual <u>CODE</u> for a certain <u>TRAIT</u>.
- There must be a gene for <u>HAIR COLOUR</u>, <u>LEFT</u>-<u>HANDEDNESS</u> and so on.
- Every <u>CHARACTERISTIC</u> we have must have a corresponding <u>GENE</u> in our chromosomes.

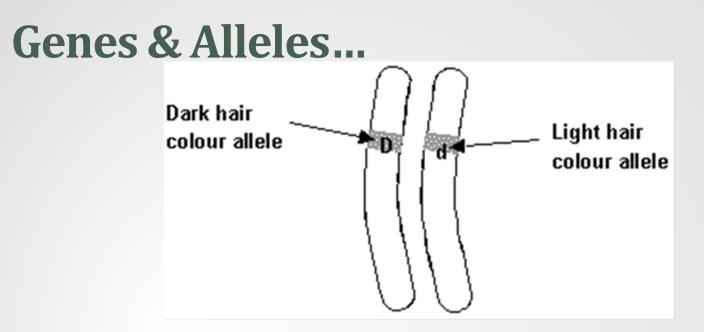


### **Genes & Alleles...**

#### <u>Alleles</u>

- An allele is a <u>FORM</u> of the <u>GENE</u>. For example in the homologous chromosomes shown in the diagram there is a <u>GENE</u> for <u>HAIR COLOUR</u>.
- Alleles for hair colour can be <u>DARK(D</u>) or <u>LIGHT(d</u>), where <u>DARK</u> hair colour is the <u>DOMINANT</u> allele.

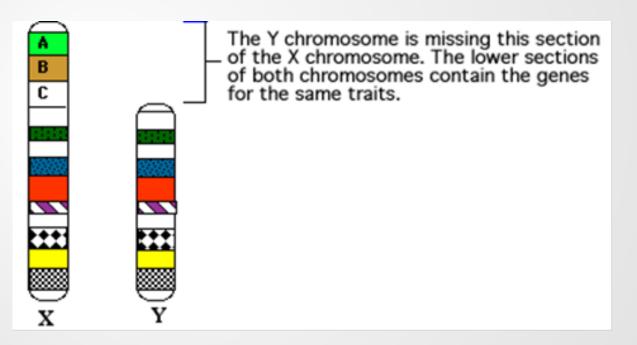




- After <u>MEIOSIS</u> has taken place, only one of the <u>HOMOLOGOUS</u> <u>CHROMOSOMES</u> will give genetic information from this parent to the offspring.
- This means the parent in the example can contribute either a <u>DARK</u> hair <u>DOMINANT</u> allele or <u>LIGHT</u> hair <u>RECESSIVE</u> allele to the offspring.
- Whether the offspring will have light or dark hair will depend on the <u>ALLELES</u> contributed by <u>BOTH</u> <u>PARENTS</u>.

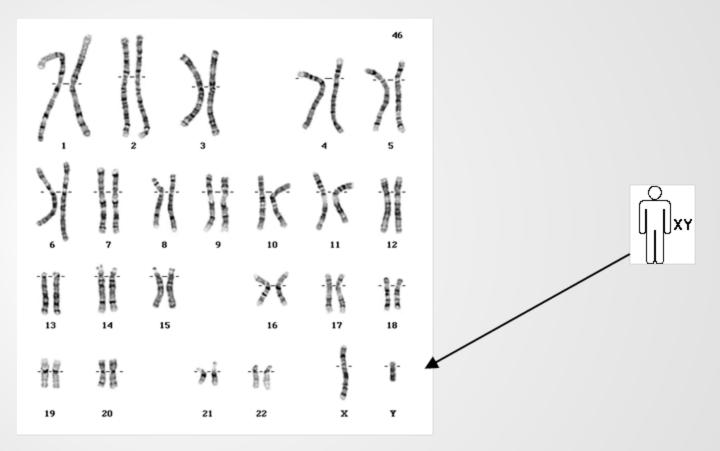
Sex-linked traits are traits carried on SEX CHROMOSOMES (X AND Y).

The male determining chromosome( $\underline{Y}$ ) has no corresponding <u>ALLELES</u> on the <u>X</u> chromosome to <u>MASK ITS EFFECTS</u>.



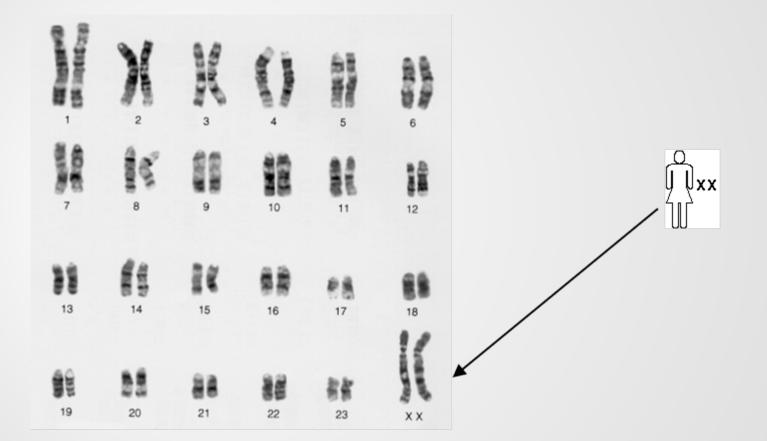
The presence of the Y chromosome causes "<u>MALENESS</u>". The female determining chromosome(<u>X</u>) does not carry <u>MALE GENES</u> of the Y chromosome

A male has XY homologous chromosomes for sex determination



The male has one X and one Y sex chromosome. Since  $\underline{Y}$  genes can't be masked by genes on the  $\underline{X CHROMOSOME}$ , he is male.

A female has XX homologous chromosomes for sex determination



The female has two <u>X</u> sex chromosomes. Since there are no male, <u>Y-BASED GENES</u>, she is female.

The X and Y chromosomes also carry genes that code for traits other than gender. Traits determined by genes on the <u>X</u> **CHROMOSOME** are called **SEX-LINKED**.

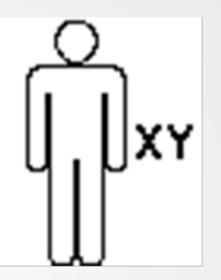
# Some of these sex-linked traits show up as **DISORDERS** like **HEMOPHILIA** and **COLOUR BLINDNESS**.

→ the genes for these disorders are <u>**RECESSIVE</u>** and found only on the <u>**X-CHROMOSOME**</u>.</u>

Ex) Colour blindness is recessive to normal vision. This is disorder mostly found in men. Why?

### Males and Sex-Linked Traits...

Males receive X chromosomes from their mothers only.



Sex-linked traits are always on the <u>X CHROMOSOME</u> and a male only has one

If he receives an X chromosome with a <u>SEX-LINKED</u> allele on it, he will always <u>DEMONSTRATE</u> that trait because there is no corresponding <u>ALLELE</u> on the <u>Y</u> chromosome to <u>MASK IT</u>.

### Females and Sex-Linked Traits...

Females receive X chromosomes from both parents and therefore can inherit sex linked traits from either parent.



If a female is to show a sex-linked trait, she must have one **DOMINANT ALLELE** on an <u>X</u> chromosome or two **RECESSIVE ALLELES** on both <u>X</u> chromosomes.

If a female receives <u>ONE RECESSIVE</u> sex-linked allele from her mother or father she <u>WILL NOT</u> show the trait, but she is a <u>CARRIER</u> and there is a probability that she will pass the sexlinked trait <u>TO ONE-HALF OF HER\_SONS</u>.

## Punnett Squares & Sex-Linked Traits ...

#### Example:

A woman who is heterozygous for colour-blindness (a carrier) has children with a man with normal vision. What genotypes and phenotypes will result?

The heterozygous mother who does not exhibit colour-blindness has a <u>50%</u> chance of producing a <u>COLOUR</u>-<u>BLIND</u> son and <u>ZERO</u> chance of producing a <u>COLOUR</u>-<u>BLIND</u> daughter.

The <u>HOMOZYGOUS DAUGHTER</u> will not have any <u>COLOUR</u>-<u>BLIND</u> offspring if she has children with a <u>NORMAL VISION</u> male. The <u>HETEROZYGOUS DAUGHTER</u> will produce offspring with the <u>SAME</u> <u>RESULTS</u> as the mother.

## Punnett Squares & Sex-Linked Traits...

#### Example 2:

What kind of offspring result from a colour-blind father and heterozygous normal mother?

The colour-blind father has

- A <u>25%</u> chance of producing a <u>COLOUR BLIND DAUGHTER</u>
- A <u>25%</u> chance of producing a <u>COLOUR-BLIND SON</u>.

If the colour-blind daughter now married a colour-blind male, the offspring would have **100% COLOUR-BLINDNESS** 

### Punnett Squares & Sex-Linked Traits...

The probability of having a homozygous colour-blind female and a colour-blind male producing offspring is not common in the general population.

If however, there were significant inbreeding, that is if relatives with a sex linked disease intermarried, then a problem can develop.

An example genetic diseases and sex linking can be seen in European aristocracy. Royalty knew no national boundaries in Europe. Intermarriage among nations was so common that a Russian Prince could have a genetic background that was mostly British. A sex-linked gene for hemophilia was introduced through marriage. Through intermarriage the female carrier would infect the males to such an extent that it became a problem.