

**Mendel** is called father of genetics. There are three laws of Mendel in respect of inheritance:

- 1. Law of dominance
- 2. Law of segregation or Law of purity of gametes or Law of splitting of hybrids.
- 3. Law of independent assortment

## 6 **(b)**

Test cross is a cross between  $F_1$  hybrid with its recessive parent.

## 7 **(a)**

Homogametic.

XY and XY type sex determination seen in many insect and mammals including humans. Males have X and Y chromosome along with autosome and females have pair of 'X' chromosome along with autosome **Parents** Phenotypes Male Female Genotypes 44A + XY 44A + XXGametes 22A + X 22A + Y22A + A22A + X22A+X 22A + XChildren 22A + X 44A + XX44 A + XYFemale 22A + Y = 44A + XY44 A + XY

Male

**Sex ratio** Female : Male = 1 : 1

**In plants** The flowering plants are mostly bisexual and lack sex chromosomes. The unisexual flowering plants tent to have XX-XY type of sex chromosomal mechanism for sex determination. The female plants are XX and male plants are XY.

### XX and XO Type of Sex Determination

Found in insect like grasshopper, cockroaches and bugs. Males have only X sex-chromosome and autosomes, female have pair of X-chromosome and autosome

Parents	Phenotypes	Male	Female		
	Genotypes	AA + XO	AA + XX		
Gametes	Α	+ X, A + O	A + X, A + Y		

### $F_1$ -generation

$$\begin{array}{c|c} A+X & A+X \\ A+X & AA+XO & AA+XO \\ A+O & AA+XO & AA+XO \\ \hline Genotypes \end{array}$$

XX-XO type of sex determination In most of cases the female produce similar sex chromosome called homomorphic. In most of cases the male produce dissimilar sex chromosome called hetermorphic

#### 8 **(a)**

In birds, usually female is designated as ZW, being heterogametic and male is designated as ZZ being homogametic.

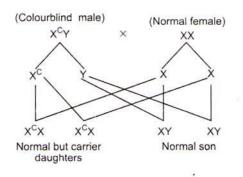
## 9 (a)

A cross of round yellow seeds (both dominant) and green wrinkled seed (both recessive) plants produced 9:3:3:1 ratio of plants (phenotypic)in  $F_2$  generation. The ratio of parental to recombinant is 10:6 here because the 9 and 1 are of parental type and 3 & 3 are recombinant.

## 10 (a)

In genetics, a test cross, first introduced by **Gregor** Johann Mendel, is used to determine weather an individual exhibiting a dominant trait is homozygous or heterozygous for that traits. More simply, test cross determines the genotype of an individual with a dominant phenotype. The test cross is defind as being a type of back cross between the recessive homozygote parents and  $F_1$ generation.

11 **(c)** 



So, all sons in the progeny will be normal.

# 12 **(d)**

When a tall pea plant (TT) is crossed with dwarf plant (tt), the  $F_1$  progeny shows all plants hybrid tall and on selfing of  $F_1$  progeny, the  $F_2$  generation

shows both tall and dwarf plant in the ratio 3 : 1. Out of three tall plants, one is pure tall (TT) and two are hybrid tall (Tt).

## 13 **(c)**

A-Two, B-Chromosomal, C-Mutation

## 14 **(c)**

A-Heterozygous, B-Unaffected, C-Carrier

### 16 **(c)**

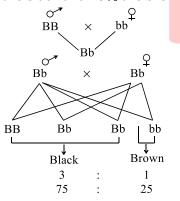
Linkage group will be equal to haploid number of chromosomes. *Pisum sativum* has seven pairs of chromosomes, therefore linkage group is also **seven**.

# 17 **(d)**

When a cross (dihybrid) is made between plants bearing round yellow (RRYY) and wrinkled green (rryy) seeds, all the plants in  $F_1$ -generation are with yellow round seeds (showing the genotype RrYy).

## 18 **(a)**

Black colour is dominant over the recessive so by cross it is easily infered that 75% of the offspring are black and 25% are brown



19 **(b)** 

There are only very few characters, which are present on the Y-chromosome of male. Like hypertrichosis. Given pedigree analysis is the example of Y-linked inheritance because all male progeny is affected

### 20 **(a)**

Haemophilia. Genetic or chromosomal symbol used for person who is having sickle-cell anaemia Ps – Hb<sup>s</sup> Hb<sup>s</sup>. Sickle-cell Anaemia (i) It is an autosome-linked recessive trait (ii) The disease is controlled by a single pair of allele Hb<sup>s</sup> and Hb<sup>s</sup> (iii) Only the homozygous individuals for Hb<sup>s</sup>, *i.e.*, Hb<sup>s</sup>Hb<sup>s</sup> show the diseased phenotype (iv) The heterozygous individuals are carriers (Hb<sup>A</sup>Hb<sup>S</sup>) (v) Due to point mutation, glutamic acid (Glu) is replaced by valine (Val) at the sixth position of  $\beta$ globin chain of haemoglobin molecule (vi) A single base substitution at sixth codon of the beta globulin gene from GAG to GUG. GAG code for glutamic acid and GUG code for valine. (vii) Hb<sup>S</sup> behaves as normal haemoglobin except under the oxygen stress where erythrocytes lose their circular shape and become sickle-shaped. As a result, the cells cannot pass through narrow capillaries. Blood capillaries are clogged and thus, affect blood supply to different organs

ANSWER-KEY										
Q.	1	2	3	4	5	6	7	8	9	10
A.	Α	В	D	В	В	B	A	A	Α	A
Q.	11	12	13	14	15	16	17	18	19	20
А.	С	D	С	С	D	С	D	Α	В	A