

Genetics :- Heredity And Variation

→ Is the branch of biological science which deals with the transmission of characteristics from one generation to the next and also the action of hereditary units called Genes as they bring about the change characteristics which they control.

→ Modern Genetics is concerned with the study of genes, the unit of heredity that control the characteristics of organisms. The term genetics was coined and used for the first time by W. Bateson (1905). Heredity or Inheritance is the characteristics transmitted from generation to generation and is therefore fixed for a particular individual.

Heredity: (i) It plays an important role in the formation of new species. Heredity involves the transfer of chromosomes from parents to offsprings or one individual to another. This indicates Chromosomal basis of heredity.

(ii) The physical basis of heredity are genes while chemical basis of heredity is DNA.

(iii) Any inherited trait such as eye colour is referred to as phenotype. All phenotypes results from the presence of a specific gene or combination of genes the genotype. In hybrid one phenotype may be dominant over another when two genotypes produces the same phenotype due to different environments one is called the phenocopy of the other since they differ in genotypes.

Pre-Mendelian views on heredity :- A number of views points were put-forward before Mendel to explain the transmission of characters from parents to offsprings. They are often called theories of blending inheritance (moist vapour theory, fluid theory, epigenesis, theory of pangenes etc.). as they believed that characters of parents blended or got mixed during their transmission to the offsprings. This theory accounted for the fact that offsprings tended to resemble their parents without all members of a population eventually averaging themselves out.

Evidence against theories of blending inheritance :- The trait of sex does not blend itself in unisexual organisms. Such an organism can either be male or female. If blending inheritance is true. The children of dark and fair coloured parents should be of intermediate colour. But intermediate colour children are often of different colours, some fair-coloured, some dark-coloured or other of intermediate colour.

2) Many individuals show ancestral characters not found in immediate parents. This phenomenon is called atavism. For ex:- (i) short tail may be found in some babies (ii) Some persons are able to move pinna or external ears.

Kolreuter (1760) a German botanist obtained fertile interspecific hybrids in tobacco. The hybrids did not resemble either of parents. Hybrids were self-pollinated. Some offsprings resembled the hybrids while remaining resembled one of the other grand parents (different characters). This proved that the traits have particular nature and remain discrete.

John Goss (1822) crossed yellow and green seeded pea varieties. The hybrids were all yellow seeded. They were self-pollinated, three types of offsprings were produced:- (i) yellow seeded (ii) green seeded (iii) with both yellow & green seeds.

Naudin (1854) concluded that on repeated crossing of hybrids the parental type appears in the offsprings showing that hybrid contains traits of both the parents though they may not be visible externally.

→ The work of the above plant breeders gave the following basic features of inheritance:-

- 1) Every character of traits has two alternative forms in which it is expressed (height \rightarrow tall / dwarf).
- 2) Traits are represented in the individuals by distinct particles which do not blend or change.
- 3) Traits may remain unexpressed for one/more generations and reappears later unchanged.
- 4) Traits may remain together in one generation and separate in a later generation.
- 5) One alternative of a trait may express more often than the other.

Terms Related to heredity \rightarrow 1) Character \rightarrow is a well defined morphological or physiological feature of an organism.

2) Traits is the distinguishing feature of a character. \rightarrow Gene \rightarrow is the inherited factor that determine the biological character of an organism.

4) Allelomorph / Allele \rightarrow a pair of contrasting characters term allele. was given by W. Bateson for alternative form of same gene e.g. T and t, Y and y, R and r are pair of allele.

5) Dominant allele \rightarrow is one of the factors of an alleles pair which can express itself whether present in homozygous or heterozygous state e.g. T (tallness in pea) R (Round seed) in pea.

6) Recessive allele \rightarrow is the factor of an allele pair which is unable to express its effect in the presence of its contrasting factor in a heterozygote e.g. t in Tt, r in Rr. The effect of recessive factor is expressed only when it is present in the pure or homozygous state. e.g. tt in dwarf pea plant.

7) wild allele \rightarrow is the one which was originally present in the population and is dominant and widespread.

8) Homozygous / pure \rightarrow organism have two similar genes/alleles for a particular characters in a homologous pair of chromosomes e.g. TT or tt.

9) Heterozygous / hybrid \rightarrow organisms containing two different alleles or individual containing both dominant and recessive genes of an allelic pair e.g. Tt is Kp heterozygous / hybrid.

10) monohybrid cross \rightarrow when only one allelic pair is considered in cross breeding.

11) Dihybrid cross \rightarrow when two allelic pairs are used in cross breeding.

12) Polyhybrid cross \rightarrow Involvement of $>$ than 2 allelic pair in cross breeding.

13) First filial generation \rightarrow 1st stage of Mendel's experiment is called first filial generation.

14) Second filial generation \rightarrow 2nd stage of Mendel's experiment. it is called second filial generation.

15) Complete penetrance \rightarrow is 100% ability of an allelic combination to produce expected phenotype. provide 100% phenotype expression ex - polydactyly, diabetes mellitus.

16) Incomplete penetrance \rightarrow is failure of an allelic combination to provide 100% phenotype expression.

17) Genotype \rightarrow is the sum total of heredity or genes make up. \rightarrow Phenotype \rightarrow is the external feature of organism.

\rightarrow Genotype remains the same throughout the life of an individual. \rightarrow Phenotype may change with the time and environment e.g. infant \rightarrow adolescent \rightarrow young \rightarrow old.

\rightarrow Genotype cannot be studied directly, it can be known through the study of ancestors mating and offspring. \rightarrow Phenotype can be known through direct observation.

\rightarrow In a given environments at time, individual with similar genotypes will produce similar phenotypes. \rightarrow Individuals with similar phenotypes may not belong to same genotype.

\rightarrow Individuals with different genotypes may have similar phenotypes e.g. tallness for TT and Tt. \rightarrow Individuals with different phenotypes not usually have different genotypes.

Mendelism! → Gregor Johann Mendel (1822-84) appropriately Ka Father of Genetics, proposed the theory of his findings on Pisum sativum in paper "Experiments on Plant hybridization" was published in the fourth volume of "Annual proceedings of Natural history society of Brunn" in 1866.

→ However, Mendel's work remained un-noticed and unappreciated for some 34 years. This is because of:-

- 1) Limited circulation of "Proceedings of Natural history society of Brunn".
- 2) Mendel's conclusions about heredity were ahead of his time.
- 3) He couldn't convince himself about his conclusion being universal since Mendel's failed to reproduce the results on Hawk-weed (Hieracium) undertaken on the suggestion of Nageli.
- 4) Lack of aggressiveness in his personality.
- 5) The scientific world was being rocked at that time by Darwin theory of evolution.

It was in 1900 that three workers independently rediscovered the principles of heredity already worked out by Mendel. They were Hugo de Vries of Holland, Carl Correns of Germany and Erich von Tschermak of Austria.

Mendel's Experiment! → Mendel conducted cross breeding experiments on garden pea (Pisum sativum). He studied the inheritance of seven different pairs of contrasting characters in this plant but considered only one pair at a time. He crossed two pea plants with alternate characters by artificial pollination. The resulting hybrids which resembled one of the two parents were then crossed with each other, both traits reappeared in the offsprings in a definite ratio of 3:1. His experiments can be depicted under the following sub-headings! →

- Selection of material! → Mendel selected garden pea as his experimental material because it had the following advantages:
- 1) The pea plants showed a number of well defined contrasting characters.
 - 2) It has perfect bisexual flowers containing both male and female parts. The flowers are predominantly self-fertile.
 - 3) Because of self-fertilization, plants are homozygous. It is therefore, easy to get pure lines for several generations.
 - 4) It was an annual plant. Its short life cycle made it possible to study several generations within a short period.
 - 5) It is easy to cultivate.
 - 6) It is easy to cross because pollen from one plant can be introduced to the stigma of another plant.

Hybridization! → Because garden pea is self-fertilizing, the anthers need to be removed before maturity. The process of removal of anthers is called emasculation.

→ At the time of cross pollination the pollen should be mature and stigma should be receptive.

→ For this cross ~~is~~ used pollen from a white flowered individual to fertilize a flower on a pea plant that produce ~~purple~~ purple flowers.

→ The stigma is protected against any pollen (from reaching it) by covering it with a bag (baggies).

→ The pollen at the dehiscence stage is brought from the plant, to be used as male parent and is directed on the stigma of emasculated flowers.

Selfing! → Flowers of F_1 generation were selfed to obtain F_2 plants. In order to avoid contamination from foreign pollens, the flowers were covered with paper bags from the beginning.
 ⇒ Mendel collected the seeds and raised a new generation of plants. ⇒ The seeds and plants raised from them constitute the second filial or F_2 generation. Further self-fertilisation produced F_3 or third filial generation.

Observation! → F_1 plants were found to have only one type of trait. F_2 generation had 2 types of plants, a large number of the traits found in F_1 generation and smaller number of the other trait not appeared in F_1 generation. In all cases the ratio was 3:1.

⇒ 1) Mendel's results can be explained using a specific example. When tall plants were crossed with dwarf plants all plants in the F_1 generation were tall. The plants used in the initial cross are referred to as P_1 and P_2 or parents.
 ⇒ 2) When the F_1 plants were self-fertilized both tall and dwarf plants were obtained in a ratio close to 3:1. Similar patterns were obtained for other six pairs of characters also.

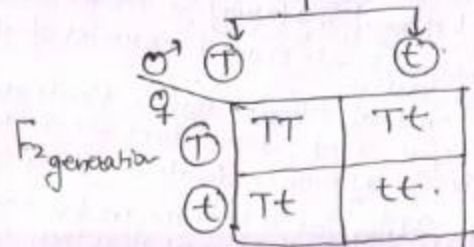
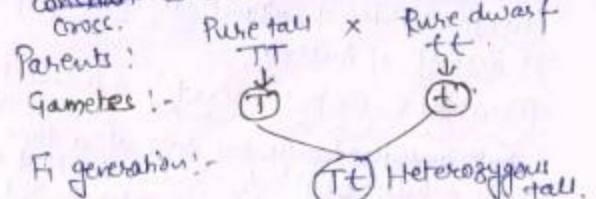
3) For any character the F_1 individuals derived from crosses between two different varieties having alternative characters showed only one of the traits and never the others. This feature was expressed as dominance of one trait over the other. The trait which appeared in the F_1 generation was called dominant and the other which did not appear in the F_1 population was called recessive.

Summary of Mendel's Experiments → Monohybrid Crosses

Character of trait studied	Parents forms crossed (P_1 cross)	F_1 products	F_2 products	Chromosome location.
1. Colour of seed	Yellow Green	All yellow	Dominant form: Yellow Recessive form: Green 3:01	1
2. Shape of seed	Round wrinkled	All Round	Round : wrinkled 2.96 : 1	7
3. Flower colour	Purple white	All-purple	Purple : white 3.15 : 1	1
4. Colour of pod	Green yellow	All green	Green : yellow 2.82 : 1	5
5. Shape of pod	Inflated Constricted	All inflated	Inflated : constricted 2.95 : 1	4
6. Position of flowers	Axial Terminal	All axial	Axial : Terminal 3.14 : 1	4
Height of plant	Tall Dwarf	All tall	Tall : Dwarf 2.84 : 1	4

Monohybrid Cross:-

The cross in which only alternate forms of a single character are taken into consideration is called monohybrid cross.



Phenotype Ratio:- 3:1
 Genotype Ratio:- 1:2:1

Dihybrid Cross!

Parents :- Yellow Round pea plant \times Green wrinkled pea plant

YYRR

yyrr

Gametes!-

YR

yr

F₁ generation!-

YyRr (Yellow Round)

Gametes of F₁

YR

Yr

yR

yr

YR YYRR

Yr YYRr

yR YyRR

yr YyRr

Yr YYRr

YYrr

yR YyRr

Yyrr

yR YyRR

YyRr

yyRR

yyRr

yr YyRr

Yyrr

yyRr

yyrr

Phenotypic Ratio :- Yellow round 9

Green Round 3

Yellow wrinkled 3

Green wrinkled 1

Summary of Mendel's Experiments And Results

Number of traits/hybrid (n)

Experiment

Type of Gametes (2ⁿ)

Number of Zygotes/offspring (gametes)²

Number of phenotypes (2ⁿ)

Number of Genotypes (3ⁿ)

Phenotypic Ratio

Genotypic Ratio

1.

Monohybrid cross (Aa Aa)

2¹ = 2

2² = 4

2¹ = 2

3¹ = 3

3:1

1:2:1

2.

Dihybrid cross (AaBb AaBb)

2² = 4

4² = 16

2² = 4

3² = 9

(3:1)² = 9:3:3:1

(1:2:1)² = 1:2:1:2:4:2:1:2:1

3.

Trihybrid cross (AaBbCc AaBbCc)

2³ = 8

8² = 64

2³ = 8

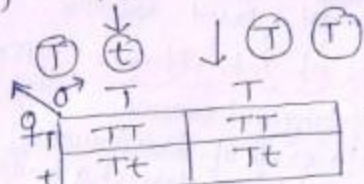
3³ = 27

(3:1)³

(1:2:1)³

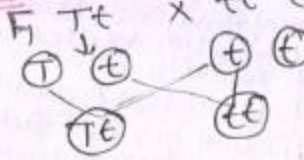
Back Cross! → A cross of F₁ hybrid with either of the two parents is K/a back cross. When F₁ offspring are crossed with the dominant parents, all the F₂ offspring develop dominant character. → On the other hand when F₁ hybrid is crossed with recessive parent, individuals with both the phenotypes appear in equal proportions while both the crosses are K/a back cross, the second one is specific as test cross.

(F₁) Tt × TT (Parental)



TT 50% pure tall : 50% Tt hybrid tall

Test Cross! → F₁ Tt × tt (Recessive parent)



Tall : Dwarf 1:1

Dihybrid Test Cross →

$P \rightarrow (YyRr) \times yyrr$ (Homozygous double recessive parent).

♂	YR	Yr	yR	yr
♀	Yr	yyRr	yyrr	yyRr
	Yellow Round	Yellow wrinkled	Green Round	Green wrinkled

Mendel's principles of inheritance →

monohybrid and polyhybrid crosses.

One gene inheritance → study of inheritance of a single pair of alleles/factors of a trait at a time (monohybrid cross) is called one gene inheritance. on the basis of his observation on monohybrid cross, Mendel proposed a set of generalisations (postulates) which resulted into the formation of following 3 laws of inheritance:—

1) Principle of paired characters → A character is represented genetically in an individual by 2 unit factors present on the same locus in the two homologous chromosomes. The 2 unit factors: two of first trait (TT), two of second trait (tt) or two of both traits (Tt).

2) Law of Dominance → It states when a pair of alleles or allelomorphs are brought together in F₁ hybrid, then only one of them expresses itself, masking the expression of other completely. In Tt F₁ hybrid:— T is expressed — so dominant is masked — so recessive.

Exceptions to principle of dominance →

1) Incomplete dominance (or partial or mosaic dominance)

Correns, 1903 — It is the phenomenon where none of the two contrasting alleles or factor is dominant. The expression of the character in a hybrid of F₁ individual is intermediate or a fine mixture of the expression of the two factors (as found in homozygous state).

→ In complete or mosaic inheritance is not an example of

Reciprocal Cross →

A set of two reciprocal crosses means that the same two parents are used in two experiments in such a way that in one experiment 'A' is used as the female parent and 'B' is used as the male parent than in the other, experiment 'A' will be used as the male parent and 'B' as the female parent. Thus the reciprocal crosses involve two crosses concerning the same characteristics but with the reversed sexes.

Mendelian principles are the rules of inheritance first discovered by Mendel. following principles or laws of inheritance based on

Pre Mendelian concept of blending inheritance because the parental types reappear in the F₂ generation. It is however considered to be an example of quantitative inheritance where only a single gene takes part involved.

2) Codominance → In it both allelic genes of a genetic trait are equally expressive i.e. the dominant character is not able to suppress the recessive characters and thus both the characters appear side by side in F₁ hybrids.

B) Law of segregation or law of purity of gametes

In F₁ the dominant phenotype appears, the recessive phenotype is not lost but reappears in F₂. This suggested that there is no blending of Mendelian factors in F₁, but they stay together and only one is expressed. At the time of the formation of gametes, these two factors obviously separate or segregate, otherwise recessive type will not appear in F₂.

→ The gametes which are formed are always pure for a particular character. A gamete may carry either the dominant or the recessive factors but not both as we find in F₁ individuals. This is why it is called either as "Principle of segregation" or as "Law of purity of gametes".

3:1 Ratio in F₂ generation of a monohybrid cross suggested that segregation of alleles does take place which can be confirmed by test cross.

Limitations of Law of Segregation → The law applies only to diploid organisms

that form haploid gamete to reproduce sexually.
→ The law only applies to traits controlled exclusively by a single gene pair in which one of the two alleles is dominant over the other.

→ The law does not apply to -

- (i) Alleles that are incompletely dominant or codominant.
 - (ii) Genes that are pleiotropic, complementary or epistatic.
- Traits caused by many gene pairs.

Inheritance of two genes →

To verify his results of monohybrid crosses, Mendel also crossed pea plants differing in two characters (dihybrid cross). This helped him to understand inheritance of two genes (i.e. two pairs of alleles) at a time.

→ It was found that inheritance of one pair of allele (one trait) does not interfere in the inheritance of other pair of allele (second trait). Based upon it, Mendel proposed a second set of generalisations (postulates) which is now called law of independent assortment.

→ Law of Independent Assortment → It states that when two pairs of independent alleles are brought together in F_1 hybrid, they show independent dominant effects. In the formation of gametes, law of segregation occurs and the factors assort independently at random.

Limitations of Law of Independent Assortment →

→ The two factors of each trait assort at random and independent of the factors of other traits at the time of meiosis (gamete genesis) and get randomly as well as independently rearranged in the offspring.

→ The law of independent assortment is applicable to only those factors or genes which are present on different chromosomes. i.e. one gene pair on one pair of homologous chromosomes and the other gene pair on another pair of homologous chromosomes. Though Mendel did not know about chromosomes, all the traits he studied were on different pairs of homologous chromosomes.

→ That is why he concluded that gene pairs segregate independently of one another. Mendel's gene pair did not show linkage.

Post Mendelism → Mendel explained inheritance in terms of discrete hereditary factors (now called genes). All patterns of inheritance could not be explained on the basis of Mendel's principle alone.

→ Bateson then proposed a hypothesis according to which more than one gene (two, three or more) may interact to produce the same character or many genes may exert influence on the development of every hereditary character with one, two or several genes having a predominant effect. Though genes are inherited as units, many interact in different patterns to produce the traits. This is called gene interaction.

Gene Interaction/Modification of F_2 ratio →

Modification of the normal phenotypic expression of genes due to interaction of their alleles and nonallelic genes. Cross correspondingly, gene interaction is of two types -

Intragenic and Intergenic

Intragenic Interaction → In it two alleles of a gene which are present on the same gene locus on the two homologous chromosomes react with each other in such a way as to produce an expression different from the normal dominant-recessive phenotype. e.g. Incomplete dominance, Codominance, multiple alleles.

Incomplete dominance → (1:2:1 ratio)

→ It is the phenomenon of neither of the two alleles being dominant so that the expression by the hybrid is intermediate between the expression of the two alleles in homozygous state. F_2 phenotypic ratio is 1:2:1 similar to genotypic ratio.

→ In Mirabilis jalapa (four o'clock) and Antirrhinum majus (snapdragon / dog flower) there are 2 types of flower colour in pure state - red and white.

→ When the 2 types of plants are crossed the hybrids are pink. When the 2 types of plants are crossed the hybrids are pink. If the latter plants of F_1 generation have pink flowers, of the latter are selfed the plants of F_2 generation are of three types - red, pink and white flowered in the ratio of 1:2:1.

→ The pink colour apparently appears either due to mixing of red and white colour (incomplete dominance) or expression of a single gene for pigmented flowers which produces only pink colour (quantitative inheritance).

Codominance (1:2:1) Ratio → may be defined as phenomenon of two alleles (different forms or Mendelian factor present on the same gene locus on homologous chromosomes) lacking dominant-recessive relationship and both expressing themselves in the organism.

⇒ Human red blood corpuscles possess two types of native antigens - M and N (Landsteiner and Levine 1927). These alleles are codominant. An individual can be MM, MN or NN, having one (MM or NN) or both the types (MN) antigens.

⇒ If a cattle with black coat is crossed with a cattle with white coat, the F₁ hybrids possess neither black, nor white coat colour but have roan colour, where black and white patches appear separately. The effect is produced due to juxtaposition of small patches of red and white colour. Hence the alleles which are able to express themselves independently when present together are called **codominant alleles**.

Multiple alleles → They are multiple forms (more than two alternatives) of Mendelian factor or gene which occur on the same gene locus distributed in different organisms in the gene pool with an organism carrying only two alleles and gametes only allele.

⇒ A well known and simplest example of multiple allelism is the inheritance of ABO blood group system in human being (discovered by Karl Landsteiner).

⇒ ABO blood group system in human being is an example of both codominant and multiple alleles. Human beings have six genotypes and four blood groups - A, B, AB and O. The blood groups are determined by two types of antigens present in the surface coating of red blood cells - A and B. The antigens occur **oligosaccharide** rich head region of a **glycoprotein**. Blood group A person have antigen A, group B have antigen B.

--- AB have both antigen while blood group O persons do not carry any antigen in the coating of their erythrocytes.

⇒ In human population 3 different alleles for ABO blood group systems are found I^A, I^B and I^O or i. I^A and I^B are mutant alleles and are dominant over I^O or i which is wild allele.

⇒ I^A and I^B are responsible for A and B antigens (glycoproteins) while I^O or i does not produce any of these A or B antigens.

⇒ A person is having only two of these three alleles and blood type can be determined by their antigen types. Six genotypes combinations are possible with three alleles.

Blood type (Phenotype)	Genotype	Antigen	Antibodies
A	I ^A I ^A / I ^A I ^O	A	b.
B	I ^B I ^B / I ^B I ^O	B	a.
AB	I ^A I ^B	Both A & B	Neither a nor b.
O	I ^O I ^O	Neither A nor B	Both a & b.

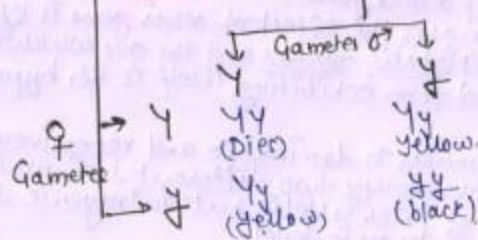
⇒ **Self-sterility in tobacco** **Nicotina** is controlled by four genes designated as S₁, S₂, S₃ and S₄.

Lethal Gene (2:1 Ratio) → They are gene which control some vital function of the organism and cause death of the organism either in homozygous recessive / homozygous dominant condition.

⇒ The individual dies in embryonic state in absolute lethality (yellow fur in mice), before reproductive maturity in sublethality (e.g. sickle-cell anaemia) and after sexual maturity in delayed lethality with the death of homozygous lethal the monohybrid ratio comes to 2:1.

⇒ **L. Cuenot** worked on the inheritance of **mouse body colour** he showed that **yellow** body colour was dominant over normal brown colour and controlled by a single gene (Y). Yellow mice could never be obtained in homozygous condition when yellow mice were crossed among themselves, segregation of yellow and brown body colour was obtained in **2:1 ratio**. ⇒ Brown individuals were pure, thus homozygous, hence dominant allele for yellow body colour is lethal in homozygous condition.

Parents: Yy (yellow) \times Yy (yellow)



Yellow 2 : 1 black

Intergenic Interaction: In intergenic or nonallelic interaction, two or more independent genes present on same or different chromosomes interact to produce a different expression e.g. epistasis, duplicate gene, Complementary genes, supplementary genes, inhibitory genes etc.

Complementary Genes (9:7 ratio)

- ⇒ Complementary genes may be defined as two or more dominant genes present on separate gene loci (non allelic pairs) which interact to produce a particular phenotypic trait, but neither of them produces the phenotypic trait in the absence of other.
- ⇒ There is complementation between two genes implying that both genes are necessary for the production of a particular phenotype.

Characteristics:
 i) Complementary genes are non-allelic genes.
 ii) They are present on separate gene loci.
 iii) Together in dominant form they produce a specific phenotypic character.

iv) Absence of either of the two is not able to produce specific phenotypic character as these genes work in collaboration.

⇒ Complementary genes were first studied by Bateson and Punnett (1906) in case of flowers colour of sweet pea (Lathyrus odoratus), here the flower colour is purple if dominant alleles of two genes C and P are present together.
 ⇒ The colour is white if the double dominant condition is absent. Bateson and Punnett crossed two white flower

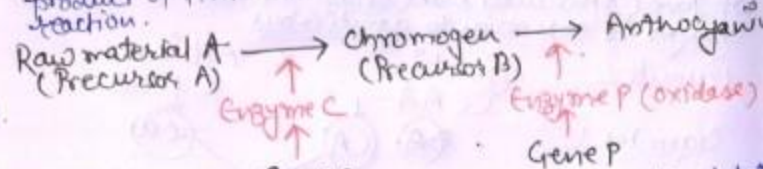
strains ($ccpp$, $ccpp$) of sweet pea and obtained purple flowered plants ($CcPp$) in the F_1 generation.
 ⇒ clearly both the parents have contributed a gene factor for the synthesis of this purple colour. The purple flowered plants of F_1 generation were then allowed to self breed.

⇒ Both purple and white flowered plants appear in the F_2 generation in the ratio of 9:7. It is the modification of the dihybrid ratio of 9:3:3:1.

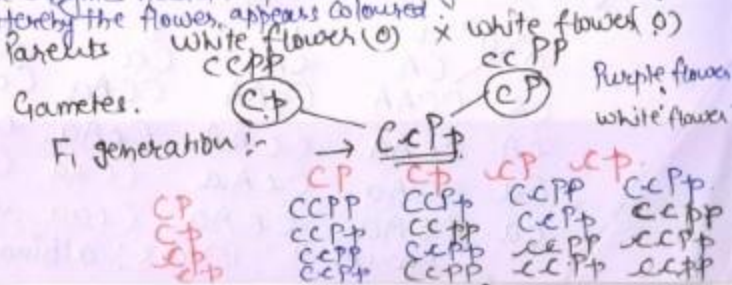
⇒ The appearance of purple colour in 9/16 population shows that the colour is determined by two dominant genes (C and P), when either of the two is absent ($ccPP$ or $CCpp$, $CcPp$, $CcPp$), the pigment does not appear.

⇒ The dominant gene C produces an enzyme which converts the raw material (precursor) into chromogen. The dominant gene P gives rise to an oxidase enzyme that changes chromogen into purple anthocyanin pigment.

⇒ This is confirmed by mixing the extract of the two types of flowers when purple colour is formed. This purple colour formation is two step reaction and the two genes cooperate to form the ultimate product. Here the end product of first reaction forms the substrate for the other reaction.



⇒ Dominance of both the genes ensures the production of both the enzymes needed to produce the pigment anthocyanin. Hence the flower appears coloured.



Supplementary genes (9:3:4 Ratio)

There are two independent genes present on different gene loci, each producing its own trait. These genes interact when present in dominant state to produce a new trait.

Characteristics! → These are a pair of non-allelic genes present on different loci on the chromosomes.

⇒ One non-allelic gene in the dominant state is able to produce its phenotypic character independently.

Parents! → Second non-allelic gene in dominant state can modify the phenotypic effect of first non-allelic gene.

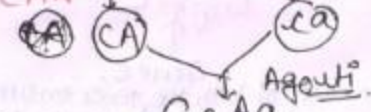
⇒ Second non-allelic gene may/may not produce its own effect independently. In collaborative supplementary genes/Supplementary genes respectively.

⇒ In mice and guinea pigs coat colour is governed by two dominant genes A and C, the agouti coloured guinea pigs have genotype CCAA. The black mice possess factor for black colour (C) but not the gene (A) for agouti colour. If gene for black colour is absent agouti is unable to express itself and mice with a genotype ccAA are albino. Here presence of gene C produced black colour and addition of gene A changes its expression to agouti colour.

Parents! -



Gametes! -



F₁ generation! -

CcAa

♀ \ ♂	CA	cA	Ca	ca
CA	CCAA	CcAA	CCaA	CcAa
cA	CcAA	ccAA	CcaA	ccAa
Ca	CCaA	CcAa	ccaa	Ccaa
ca	CcAa	ccAa	Ccaa	ccaa

Agouti 9! Black 3! albino 4.

Epistasis (12:3:1 ratio)!

Epistasis is the interaction between genes present at two separate loci in which one gene suppresses or masks the expression of other gene.

⇒ The gene that masks the effect of other gene is K_h inhibiting or **epistatic factor** and the one which is being prevented from exhibiting itself is K_h hypostatic **factor**.

⇒ Although it is similar to dominance and recessiveness but the two factors occupy two different loci. While dominance works at intra-allelic but intragenic level, epistasis works at inter-genic level.

⇒ Epistasis reduces the number of phenotypes appearing in F₂ generation of dihybrid cross. Thus instead of normal 9:3:3:1 dihybrid ratio, epistasis may result in the ratio of 9:3:4. (recessive epistasis) 12:3:1/13:3 (dominant epistasis).

Dominant epistasis! → A dominant epistatic allele suppresses the expression of a non-allelic gene whether the latter is dominant or recessive.

⇒ Fruit colour of summer squash (*Cucurbita pepo*) is governed by a gene which produces yellow colour in dominant state (Y) and green colour in recessive state (yy). There is a non-pigment forming epistatic gene which in dominant state (W-) produces white colour (W-; Y-; W-; yy) while in recessive state it allows the yellow (wwY-) and green (wwyy) expression.

Parents! - WWYY (white fruit) × wwyy (green fruit).

Gametes! -

WY wY WY wY

	WY	wY	WY	wY
WY	WWYY	WwYy	WWYY	WwYy
wY	WwYy	wwYY	WwYy	wwYY
WY	WWYY	WwYy	WWYY	WwYy
wY	WwYy	wwYY	WwYy	wwYY