Chapter 01

Principles of Inheritance and Variation





TOPIC WISE QUESTIONS



MENDELISM

- **Q.1** F_1 progeny of Mendelian dihybrid cross produces:
 - (1) Two types of pollen grains
 - (2) Four genotypes of gametes
 - (3) Two types of eggs
 - (4) Four types of pollens only
- Q.2 When Mendel self hybridised the F₁ plants (RrYy), he found that dominant and recessive traits of one character are segregated in:
 - (1) 9:1 ratio
- (2) 3 : 3 ratio
- (3) 10 : 6 ratio
- (4) 3: 1 ratio
- Q.3 Mendel published his work on inheritance of characters in 1865 but it remained unrecognized till 1900 because:
 - (a) He could not provide any physical proof for the existence of factors
 - (b) His concept of factors as stable, discrete units that controlled the expression of traits did not find acceptance from the contemporaries
 - (c) Mendel's approach of using mathematics to explain biological phenomenon was totally old
 - (d) Communication was not easy (as it is now)
 - (1) (a), (b) and (c) are correct
 - (2) (c) and (d) are correct
 - (3) (a), (b) and (d) are correct
 - (4) Only (a) is correct
- Q.4 Heterozygous tall and violet flowered pea plants were selfed and total 512 seeds are collected. What will be total number of seeds for both heterozygous traits?
 - (1) 128
- (2)256
- (3) 384
- (4) 64
- **Q.5** Mark the odd one (with respect to F_2 generation of Mendelian dihybrid cross):
 - (1) Frequency of TtRR genotype = 12.5%
 - (2) Frequency of ttrr genotype = 6.25%
 - (3) Frequency of TTRR genotype = 6.25%
 - (4) Frequency of ttRr genotype = 25%
- **Q.6** Mendel studied seven pair of contrasting traits in Pea plant. Which is correct option in given alternatives?

- (1) 2 characters flower based, 2 colour based, 3 seed based, 1 height based
- (2) 2 characters flower based, 3 colour based, 1 height based, 2 seed based, 2 pod based
- (3) 3 colour based, 1 flower based, 3 height based
- (4) 3 Characters pod based, 2 stem based, 1 height based, 3 colour based
- **Q.7** Select incorrect one (w.r.t. reciprocal cross):
 - (1) To know whether the alleles are present on sex chromosomes or autosomes.
 - (2) It is made to eliminate to effect of nuclear trait.
 - (3) Two individuals with contrast genotypes are involved.
 - (4) Results are not changed for autosomal traits.
- Q.8 Mendel conducted hybridisation experiments on garden peas between
 - (1) 1756 1763
- (2) 1856 1863
- (3) 1656 1663
- (4) 1956 1963
- Q.9 Which of the following combination seems to have some linkage in character selected by Mendel?
 - (1) Stem height and pod colour
 - (2) Flower colour and flower position
 - (3) Seed shape and seed colour
 - (4) Plant height and pod shape
- Q.10 A diploid organism is heterozygous for five loci and homozygous for 2 loci, how many types of gametes can be produced?
 - (1) 128
- (2) 32
- (3)
- (4) 14
- **Q.11** In F₂ generation of a Mendelian dihybrid cross (TTRR x ttrr):
 - (1) Tall plants and violet flowered plants are obtained in 1 : 1 frequency
 - (2) Ratio of parental and non-parental plants is 1:15
 - (3) Recombinant plants are obtained in 1 : 1 frequency
 - (4) More than one option is correct



PRINCIPLES OF INHERITANCE AND VARIATION

- **Q.12** When a heterozygous tall pea plant of F_1 generation upon self fertilization produces tall and dwarf phenotypes. It proves the principle of:
 - (1) Dominance
 - (2) Segregation
 - (3) Independent assortment
 - (4) Inheritance & purity of gametes
- Q.13 A cross between AaBB X aaBB yields a genotypic ratio of:
 - (1) 1 AaBB: 1 aaBB
 - (2) 1 AaBB: 3 aaBB
 - (3) 3 AaBB: 1 aa BB
 - (4) All AaBb
- **Q.14** How many types & in what ratio the gametes are produced by a dihybrid heterozygous:
 - (1) 4 types in the ratio of 9:3:3:1
 - (2) 2 types in the ratio of 3:1
 - (3) 3 types in the ratio of 1:2:1
 - (4) 4 types in the ratio of 1:1:1:1
- **Q.15** How many gametes are produced in F_1 generation of a trihybrid:
 - (1) 3
- (2) 4
- (3) 8
- (4) 16
- Q.16 Which genotype represents a true dihybrid condition:
 - (2) Tt rr (3) Tt Rr (1) tt rr (4) TT Rr
- Q.17 In a cross between a pure tall plant with green pod & a pure short plant with yellow pod. How many short plants are produced in F₂ generation out of 16:
 - (1) 1
- (2) 3
- (3)4
- (4)9
- **Q.18** In a dihybrid cross between AABB and aabb the ratio of AABB, AABb, aaBb, aabb in F₂ generation is:
 - (1) 9:3:3:1
- (2) 1:1:1:1
- (3) 1:2:2:1
- (4) 1:1:2:2
- Q.19 Crossing AABB and aabb, the probability of AaBb would be in F_2 generation:

 - (1) 1/16 (2) 2/16 (3) 8/16
- (4) 4/16
- **O.20** If 120 Plants of Antirrhinum plants are produced on crossing pure red and pure white flowered plants, than the ratio of offsprings will be:
 - (1) 90 Red: 30 White flowered
 - (2) 30 Red: 90 White flowered
 - (3) 60 pink: 60 White flowered
 - (4) All pink flowered
- **Q.21** An individual with two identical members of a pair of genetic factors is called:
 - (1) Heteromorphic
- (2) Heterozygote
- (3) Homomorphic
- (4) Homozygote

- **Q.22** Two allelic genes are located on:
 - (1) The same chromosome
 - (2) Two homologous chromosomes
 - (3) Two non-homologous chromosomes
 - (4) Any two chromosomes
- Q.23 Mendel's law of segregation is based on separation of alleles during:
 - (1) Gamete formation
 - (2) Seed formation
 - (3) Pollination
 - (4) Embryonic development
- **Q.24** Which technique is used by Mendel for hybridisation:
 - (1) Emasculation
- (2) Bagging
- (3) Protoplast fusion (4) 1 & 2 both
- **Q.25** When flowers are unisexual then emasculation is done in:
 - (1) Female
- (2) Male
- (3) 1 and 2 both
- (4) None of these
- Q.26 How many plants are dihybrid in F₂ generation of dihybrid cross:
 - (1) One
- (2) Two (3) Four (4) Sixteen
- **0.27** When a plant have two alleles of contrasting characters, it is called:
 - (1) Homozygous
- (2) Dioecious
- (3) Heterozygous
- (4) Monoecious
- **Q.28** From a single ear of corn, a farmer planted 200 kernels which produced 140 tall & 40 short plants. The genotypes of these offsprings are most likely:
 - (1) TT, Tt and tt
- (2) TT and tt
- (3) TT and Tt
- (4) Tt and tt
- **Q.29** A useful process for determining whether an individual is homozygous or heterozygous is:
 - (1) Cross-breeding
- (2) Self fertilization
- (3) Back crossing
- (4) Test cross
- Q.30 Heterozygous tall plants were crossed with dwarf plants. What will be the ratio of dwarf plants in the progeny:
 - (1)50%
- (2)25%
- (3)75%(4) 100%
- Q.31 If the cell of an organism is heterozygous for two pairs of genes represented by Aa, Bb, undergoes meiosis, then the possible genotypic combination of gametes will be:
 - (1) AB, Ab, aB, ab
- (2) AB, ab
- (3) Aa, Bb
- (4) A, a, B, b
- **Q.32** Allele is the :
 - (1) Alternative form of gene pair
 - (2) Total number of genes for a trait
 - (3) Total number of chromosomes of a haploid set



- (4) Total number of genes present on a chromosome
- **Q.33** Genetic constitution of an individual represented by :
 - (1) Genome
- (2) Genotype
- (3) Phenotype
- (4) Karyotype
- **Q.34** Genes do not occur in pairs in:
 - (1) Zygote
- (2) Somatic cell
- (3) Endosperm cell
- (4) Gamete
- **Q.35** According to Mendelism, which character is showing dominance:
 - (1) Terminal position of flower
 - (2) Green colour in seed coat
 - (3) Wrinkled seeds
 - (4) Green pod colour
- **Q.36** A gene said to be dominant if:
 - (1) It express its effect only in homozygous stage.
 - (2) It is expressed only in heterozygous condition
 - (3) It is expressed both in homozygous and heterozygous condition.
 - (4) It is never expressed in any condition.
- Q.37 Mendel's principle of segregation means that the germ cells always receive
 - (1) Either one allele of father or one allele of mother
 - (2) One pair to alleles
 - (3) One quarter of the genes
 - (4) Any pair of alleles
- Q.38 If a homozygous tall plant is crossed with a dwarf plant, what shall be the ratio of plants in off springs:
 - (1) All heterozygous tall
 - (2) Two tall and Two dwarf
 - (3) 1:2:1
 - (4) All homozygous dwarf
- **Q.39** The process of mating between closely related individuals is:
 - (1) Out-breeding
- (2) Inbreeding
- (3) Hybridisation
- (4) Heterosis
- **Q.40** Marriages between close relatives should be avoided because it includes more :
 - (1) Recessive alleles to come together
 - (2) Mutations
 - (3) Multiple births
 - (4) Blood group abnormalities
- **Q.41** An inherited character and its detectable variant is termed as:
 - (1) Unit factor
- (2) Trait
- (3) Genetic profile
- (4) Genotypic character

- **Q.42** When an F_1 individual is crossed with its either of the two parent. Then it is known as :
 - (1) Test cross
- (2) Back cross
- (3) Reciprocal cross (4) Monohybrid cross
- **Q.43** How many types of genotypes are formed in F_2 progeny obtained from self pollination of a dihybrid F_1 :
 - (1)9
- (2) 3
- (3)6
- (4) 1
- **Q.44** If a dwarf plant is treated with gibberellins it becomes tall and this plant now crosses with pure tall plant then progeny of first generation (F_1) is:
 - (1) All dwarf
 - (2) All tall
 - (3) 75% tall and 25% dwarf
 - (4) 75% dwarf and 25% tall
- Q.45 Which of the following is correct for test-cross
 - (1) $TT \times Tt$
- (2) $tt \times TT$
- (3) $Tt \times tt$ (4) $tt \times tt$
- **Q.46** If a cross is made between AA and aa, the nature of F_1 progeny will be:
 - (1) Genotypically AA, phenotypically a
 - (2) Genotypically Aa, phenotypically a
 - (3) Genotypically Aa, phenotypically A
 - (4) Genotypically aa, phenotypically A
- Q.47 A pure tall and a pure dwarf plant were crossed to produce off springs. Off springs were self crossed, then find out the ratio between true breeding tall to true breeding dwarf:
 - (1) 1:1
- (2) 3:1
- (3) 2:1
- (4) 1:2:1
- Q.48 If hybrid red flowered plants of pea are crossed back to pure red flowered parent, the progeny will show:
 - (1) All red flowered plants
 - (2) White flowered plants
 - (3) 50% red and 50% white flowered plants
 - (4) 3 Red: 1 white flowered plants
- Q.49 A cross between pure tall pea plant with green pods and dwarf pea with yellow pods will produce tall F_2 plants, out of 16:
 - (1) 15
- (2) 13
- (3) 12
- (4) 7
- **Q.50** Mendel's Principle of segregation means that the germ cells always receive:
 - (1) One pair of alleles
 - (2) One quarter of the genes
 - (3) One of the paired alleles
 - (4) Any pair of alleles



Q.66 Incomplete dominance occurs in :

(2) Antirrhinum

(1) Mirabilis

Q.51 How many types of genetically different gametes Q.60 Mark the correct statement (with respect to sickle will be produced by a heterozygous plant having cell anaemia): (1) Homozygous individuals for HbSHbS are the genotype AABbCc? apparently unaffected (1) Two (2) Four (3) Six (4) Nine (2) Heterozygous individuals (Hb^AHb^S) appear **Q.52** Mendel observed that all the F_1 progeny plants: apparently unaffected (1) Resembled either one of the parents (3) Heterozygous individuals are affected as well (2) Resembled neither of the parents as carrier (3) Resembled both of the parents (4) Homozygous individuals for Hb^A show the (4) Shows 3: 1 ratio diseased phenotype **Q.53** According to Mendel, "factors" or "genes": **Q.61** The defect sickle-cell anaemia is caused by the (1) Are the units of inheritance _ of glutamic acid by valine at the 6th (2) Contain information that is required to position of the globin chain of the express a particular trait haemoglobin molecule: (3) Both 1 and 2 (1) Substitution, β (2) Deletion, α (4) Do not contain information required to (3) Duplication, β (4) Translocation,α express a particular trait. **Q.54** The segregation of alleles is a random process **Q.62** In incomplete dominance: and so there is a _____ chance of a gametes (1) Dominant trait is completely expressed in F_1 containing either allele: generation (1)25%(2) 50%(2) Phenotypic and genotypic ratio are different (4) 100% (3)75%(3) Two dominant alleles are needed to express Q.55 The phenotype of any character will not be the complete dominant trait affected if the modified allele produces: (4) F₁ individuals have the equal traits of both (1) Normal enzyme parents (2) Non-functional enzyme Q.63 Progeny with blood group 'O' can not be obtained (3) No-enzyme at all (4) 2 and 3 both in cross: (1) A x A(2) A x B **Q.56** Genes which code for a pair of contrasting traits (3) O x AB (4) B x B are known as :-(1) Alleles (2) Non alleles **Q.64** In *Mirabilis* and *Antirrhinum* plant the (3) Pseudoalleles (4) Isoalleles appearance of the pink hybrid (Rr) between cross of a red (RR) and white (rr) flower parent **Q.57** The recessive characters are: indicates: (1) Only expressed in heterozygous condition (2) Only expressed in homozygous condition (1) Incomplete dominance (3) Do not blend in heterozygous condition (2) Segregation (4) Always impure (3) Dominance (4) Heterosis **GENE INTERACTIONS Q.58** In garden pea, starch is synthesised effectively in: Q.65 In Mirabilis jalapa when homozygous red (1) Heterozygous round seeded plants flowered and white flowered plants are crossed, (2) Homozygous round seeded plants all F₁ plants have pink coloured flowers. In F₂ (3) Wrinkled seeded plants produced by selfing of F₁ plants, red, pink, white (4) Pure and hybrid round seeded plants flowered plants would appear respectively in the Q.59 When a pink flowered Antirrhinum plant is test ratio of: crossed, then phenotypic ratio in resulting (1) 1:1:2 (2) 2:1:1 (3) 1:0:1 (4) 1:2:1 progenies is:



(1) 1 Red: 1 White

(2) 3 Red: 1 White

(3) 2 Pink: 1 White (4) 1 Pink: 1 White

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	(3) Andulasion fowl (4) All of the above		pollinated to obtain a phenotypic ratio of 1:2:1, it						
Q.67	Which cross yields red, white and pink flowers variety of dog flower: (1) RR X Rr (2) Rr X RR (3) Rr X Rr (4) Rr X rr		has to be a case of: (1) Incomplete dominance (2) Dominance (3) Recessive epistasis (4) Pleiotropic effect of genes						
Q.68	What shall be ratio in offspring when a roan cow is crossed with a white bull: (1) 1:2:1 (2) 3:1 (3) 1:1 (4) All roan	Q.76	A gene that shows its effect on more than one character is: (1) Polygene (2) Pleiotropic gene (3) Multifactor gene (4) Multiple gene						
Q.69	In a dihybrid cross, when one pair of alleles show incomplete dominance, genotypic ratio comes to: (1) 3:6:3:1:2:1 (2) 1:2:2:4:1:2:1:2:1 (3) 9:3:3:1	Q.77 Q.78	In multiple allele system a gamete possesses: (1) Two alleles (2) Three alleles (3) One allele (4) Several alleles Blood grouping in humans is controlled by: (1) 4 alleles in which I ^A is dominant (2) 3 alleles in which I ^A and I ^B are dominant						
	(4) 1 : 2 : 1		(3) 2 alleles in which none is dominant						
	 Which of the following conditions represent a case of co-dominant genes: (1) A gene expresses itself, suppressing the phenotypic effect of its alleles (2) Genes that are similar in phenotypic effect when present separately, but when together interact to produce a different trait (3) Allele, both of which interact to produce a trait, which may resemble either of the parental type. (4) Alleles, each of which produces an independent effect in heterozygous condition. ABO blood group is an example of: (1) Epistasis 		 (4) 3 alleles in which I^A is recessive 9 Multiple alleles are present: (1) In different chromosomes (2) At different loci on chromosome (3) At the same locus on homologo chromosomes (4) At the non homologous chromosome (4) At the non homologous chromosome (5) Pleiotropic genes have multiple phenotype effect. (6) Multiple alleles exhibit same phenotype expression. (7) Polygenes exhibit continuous variation. 						
	(2) Multiple allelism (3) Pleiotropism (4) Complementary genes	arni	 Statement (A), (B) and (C) are correct Statement (A), (C) correct and (B) is incorrect Statement (A), (B) and (C) are incorrect 						
Q.72	A child has blood group 'O'. His parents blood group cannot be: (1) B and O (3) AB (2) A and O (4) A and B	Q.81	 (4) Statement (B) and (C) are correct and (A) is incorrect Sickle cell anaemia induces due to: (1) Change of Amino Acid in α – chain of 						
	If one parent has blood group A and the other parent has blood group B. The offsprings have which blood group: (1) AB only (2) O only (3) B only (4) A, B, AB, O		 Haemoglobin (2) Change of Amino Acid in β – chain of Haemoglobin (3) Change of Amino Acid in both α and β chain of Haemoglobin 						
Q.74	A child of O blood group, has B-blood group father, the genotype of father would be : $(1)\ I^0I^0 \qquad (2)\ I^BI^B \qquad (3)\ I^AI^B \qquad (4)\ I^BI^0$	6.65	(4) Change of Amino acid either α or β chain of Haemoglobin						
Q.75	When a red flowered plant was cross pollinated by white flowered one and the offspring were self	Q.82	Incomplete dominance is absent in: (1) Pisum sativum (2) Antrrhinum majus						



- (3) Human ABO blood groups
- (4) Both Pisum sativum and Antirrhinum majus
- Q.83 A man with blood group B marries a female with blood group A and their first child is having blood group B. What is the genotype of child:
 - (1) I^AI^B
- $(2) I^{A}I^{O}$
- (3) I^BI^O
- (4) I^BI^B
- **Q.84** A child with mother of blood group A and father of blood group AB, will not have which of the following blood group:
 - (1) A
- (2) B
- (3) AB
- (4) O
- **Q.85** If mother has blood group B, father has A group, the offspring will be of:
 - (1) A
- (2) O
- (3) AB
- (4) Any of the above
- Q.86 Sickel cell anemia is the result of mutation in the haemoglobin gene:
 - (1) Frame shift
- (2) Deletion
- (3) Point
- (4) None of the above
- **Q.87** In the inheritance of flower colour in dog flower plant, the F_1 has a phenotype that:
 - (1) Resembles both of the parents
 - (2) Does not resemble either of the two parents
 - (3) Resembles with only one parent
 - (4) 1 and 3 both
- **Q.88** The three different alleles of human ABO blood types will produce how many genotypes & phenotypes respectively:
 - (1) 4 and 6
- (2) 6 and 4
- (3) 6 and 6
- (4) 4 and 4
- **0.89** Find out the correct match:
 - (1) F_1 resembled either of the parents Dominance
 - (2) F₁ resembled in between -incomplete dominance
 - (3) F₁ resembled both parent Co-dominance
 - (4) All are correct
- Q.90 Which of the following condition is true for codominance:
 - (1) Phenotype of F_1 resembled either of the two parents
 - (2) Phenotype of F_1 did not resemble either of two parents
 - (3) Phenotype of F_1 resembles both parents
 - (4) More than one option is correct
- **0.91** In sickle cell anaemia:
 - mutant haemoglobin undergoes polymerisation under low oxygen

- tension causing the change in the shape of **RBC**
- (2) Substitution of Glutamic acid by valine at the sixth position of the α -chain of haemoglobin
- (3) The mutant haemoglobin undergoes polymerization under high oxygen tension causing the change in shape of RBC
- (4) α-globin chain is modified
- **Q.92** In case of ABO blood group allele I^A and I^B if present together then:
 - (1) Ony I^A allele expresses
 - (2) Only I^B allele expresses
 - (3) Both I^A and I^B alleles express
 - (4) None of these

POLYGENIC INHERITANCE

- **Q.93** A polygenic inheritance in human beings is:
 - (1) Skin colour
 - (2) Sickle cell anaemia
 - (3) Colour blindness
 - (4) Phenylketonuria
- Q.94 Inheritance of height in human beings is an example of:
 - (1) Complementary gene
 - (2) Monogenic inheritance
 - (3) Polygenic inheritance
 - (4) Mendelian inheritance
- **Q.95** In polygenic inheritance trait which controlled by three pairs of genes. Two individuals which are heterozygous for three alleles, crossed each other. Such type of cross produces the phenotypic ratio:
 - (1) 1:2:1
- (2) 9:3:3:1
- (3) 1:4:6:4:1
- (4) 1:6:15:20:15.:6:1
- **Q.96** A polygenic trait is controlled by 3 genes A, B and C. In a cross AaBbCc x AaBbCc, the phenotypic ratio of the offsprings was observed as:

1:6:x:20:x:6:1

What is the possible value of x?

(1) 3

(2)9

(3) 15

(4) 25

CHROMOSOMAL THEORY OF INHERITANCE, LINKAGE, SEX LINKAGE & RECOMBINATION

- Q.97 Which of the following statement chromosomal theory of inheritance is incorrect?
 - (1) Pairing and separation of a pair of chromosomes would lead to the segregation of a factor they carried
 - (2) Behaviour of chromosomes is identical to the behaviour of genes
 - (3) The two alleles of a gene pair are located on homologous sites on homologous chromosomes
 - (4) Chromosomes as well as genes occur in pairs



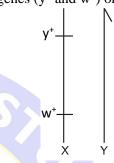
- **Q.98** Fruit flies are one of the best materials for genetic studies because of all, except:
 - (1) Ability to grow on simple synthetic medium in the laboratory
 - (2) Short life span
 - (3) Production of a large number of progeny in each mating
 - (4) Presence of few externally visible and identifiable contrasting traits
- **Q.99** Generation of non-parental gene combinations is termed as:
 - (1) Linkage
- (2) Polyploidy
- (3) Recombination
- (4) Aneuploidy
- **Q.100** Which of the following phenomena leads to variation in DNA?
 - (1) Linkage, mutation
 - (2) Recombination, linkage
 - (3) Mutation, recombination
 - (4) Aneuploidy, linkage
- **Q.101** Which of the following trait shows transmission from carrier female to male progeny?
 - (1) Autosomal dominant
 - (2) X-linked recessive
 - (3) Y-linked recessive
 - (4) X-linked dominant
- Q.102_(A)__used the frequency of recombination between gene pairs on the __(B)__ as a measure of the distance between genes and mapped their position on the chromosome.

(A)

(B)

- (1) Morgan
- Same chromosome
- (2) Sturtevant
- Different chromosomes
- (3) Morgan
- Different chromosomes
- (4) Sturtevant
- Same chromosome
- Q.103In which of the following disorder a single protein that is a part of the cascade of proteins involved in blood clotting is affected?
 - (1) Thalassemia
- (2) Sickle-cell anaemia
- (3) Haemophilia
- (4) Phenylketonuria
- Q.104A normal woman, whose father had colour blindness, married a normal man. What is the chance of occurrence of colour blindness in the progeny?
 - (1) 25%
- (2) 50%
- (3) 100% (4) 75%
- Q.105The chromosome maps are not accurate maps because
 - (1) Crossing over frequency is higher than recombination frequency

- (2) One crossing over interferes and increases the frequency of nearby crossing over
- (3) Crossing over frequency decreases towards the ends of chromosome
- (4) Heterochromatin increases crossing over
- **Q.106**How many types of gametes will be produced by a *Drosophila* having following arrangement of two genes (y⁺ and w⁺) on X-chromosome?



(1) 2

(2)4

(3) 1

(4) 8

Q.107 What is the inheritance of colour blindness of offsprings where both parents having a normal vision but mother has a recessive gene for colourblindness:

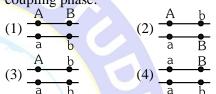
Son	Daughter
(1) 50%	Nil
(2) 100%	Nil
(3) Nil	100%
(4) Nil	Nil

- Q.108 What would be the nature of children if a colour blind woman marries a normal man:
 - (1) Colour blind daughter & normal sons
 - (2) Colour blind sons and carrier daughters
 - (3) Normal sons & carrier daughters
 - (4) Normal sons & Normal daughters
- Q.109A colourblind daughter is born when:
 - (1) Father is colour blind, mother is normal
 - (2) Mother is colour blind, father is normal
 - (3) Mother is carrier, father is normal
 - (4) Mother is carrier, father is colour blind
- **Q.110**Which of the following is not a sex linked characters:
 - (1) Haemophilia
- (2) Colour blindness
- (3) Hypertrichosis
- (4) Baldness
- Q.111A woman with normal vision marries a man with normal vision and gives birth to a colourblind son. Her husband dies and she marries a colourblind man. What is the probability of her children having the abnormality?



- (1) 50% colourblind sons + 50% colourblind daughters
- (2) All sons colourblind and daughter carrier
- (3) All daughter colourblind and sons normal
- (4) 50% sons colour blind and all daughters normal
- Q.112In a cross between individuals homozygous for (a, b) and wild type (+ +). In this cross 700 out of 1000 individuals were of parental type. Then the distance between a and b is:
 - (1) 70 map unit
- (2) 35 map unit
- (3) 30 map unit
- (4) 15 map unit
- Q.113 What ratio is expected in offsprings if father is colour blind and mother's father was colour blind:
 - (1) 50% daughter colour blind
 - (2) All the sons are colour blind
 - (3) All the daughters colour blind
 - (4) All the sons are normal
- Q.114 There are three genes a, b, c percentage of crossing over between a and b is 20%, b and c is 28% and a and c is 8%. What is the sequence of genes on chromosome:
 - (1) b, a, c (2) a, b, c (3) a, c, b (4) b, c, a
- Q.115 When a cluster of genes show linkage behaviour they:
 - (1) Do not show a chromosome map
 - (2) Show recombination during meiosis
 - (3) Do not show independent assortment
 - (4) Induce cell division
- **Q.116**Genetic Map is one that:
 - (1) Establishes sites of the genes on a chromosome
 - (2) Establishes the various stages in gene evolution
 - (3) Shows the stages during the cell division
 - (4) Shows the distribution of various species in a region
- Q.117One of the genes present exclusively on the X-chromosome in humans is concerned with:
 - (1) Baldness
 - (2) Red green colour blindness.
 - (3) Facial hair/Moustaches in males.
 - (4) Night blindness.
- **Q.118**The recessive genes located on X–chromosome in humans are always:
 - (1) Expressed in females
 - (2) Lethal
 - (3) Sub-lethal
 - (4) Expressed in males

- Q.119A colourblind man marries with normal eyed woman where there is no historical record of colourblindness. The first offspring of this couple is daughter. Find out the possibility of appearance of colourblindness of this daughter?
 - (1) 50% (2) 25% (3) 0%
 - 3) 0% (4) 75%
- **Q.120**If father shows normal genotype and mother shows a carrier trait for hemophilia:
 - (1) All the female children will be carrier
 - (2) A male child has 50% chances of active disease
 - (3) Female child has probability of 50% to active disease
 - (4) All the female children will be colourblind
- **Q.121**Which of the following show linkage group in coupling phase:



- Q.122 Which statement is incorrect about linkage:
 - (1) It helps in maintaining the valuable traits of new varieties
 - (2) It helps in forming new recombinants
 - (3) Knowledge of linkage helps the breeder to combine all desirable traits in a single variety.
 - (4) It helps in locating genes on chromosome
- Q.123 A woman with normal vision, but whose father was colour blind, marries a colour blind man. Suppose that the fourth child of this couple was a boy. This boy:
 - (1) Must have normal colour vision
 - (2) May be colour blind or may be normal vision
 - (3) Will be partially colour blind since he is heterozygous for the colour blind mutant allele
 - (4) Must be colour blind
- Q.124Haemophilia is more commonly seen in human males than in human females because:
 - (1) This disease is due to a Y-linked recessive mutation
 - (2) This disease is due to an X-linked recessive mutation
 - (3) This disease is due to an X-linked dominant mutation
 - (4) A greater proportion of girls die in infancy
- Q.125 If Mendel has chosen to study traits determined by linked genes he would not have discovered:
 - (1) Law of segregation



- (2) Law of dominance
- (3) Law of independent assortment
- (4) Law of unit character
- **Q.126**The first attempt to show linkage in plants was done in :
 - (1) Pisum sativum
 - (2) Lathyrus odoratus
 - (3) Zea mays
 - (4) Oenothera lamarckiana
- **Q.127** If there were only parental combinations in F₂ of a dihybrid cross then Mendel might have discovered:
 - (1) Independent assortment
 - (2) Atavism
 - (3) Linkage
 - (4) Repulsion
- Q.128Chromosomal theory of inheritance was proposed by
 - (1) Sutton & Boveri (2) Sturtevant
 - (3) Morgan
- (4) Henking
- Q.129 If distance between gene on chromosome is more, then gene shows:
 - (1) Weak linkage
- (2) Strong linkage
- (3) Less crossing
- (4) 1 and 3 both
- Q.130Linked gene shows:
 - (1) Always parental combination
 - (2) Sometimes new combinations
 - (3) Always new combination
 - (4) New combination more
- Q.131The number of linkage groups in a cell having 10 pairs of chromosomes are:
 - (1) 5
- (2) 10
- (3) 15
- (4) 20
- Q.132The association of parental characters combinations in the offsprings of a dihybrid is in excess to non-parental combinations is said to be due to:
 - (1) Co-dominance
- (2) Blending inheritance
- (3) Linkage
- (4) Duplicate genes
- **Q.133** A phenomenon which works opposite to the linkage is:
 - (1) Independent assortment
 - (2) Crossing-over
 - (3) Segregation
 - (4) Mutation
- **Q.134**Cross over value (COV) of gene A and B is 5% while COV of genes B and C is 15% the possible sequence of these genes on chromosome is :
 - (1) A-B-C
- (2) C-A-B
- (3) B-C-A
- (4) Both (1) and (2)

- Q.135 A diseased man marries a normal woman. They get three daughters and five sons. All the daughters were diseased and sons were normal. The gene of this disease is:
 - (1) Sex linked dominant
 - (2) Sex linked recessive
 - (3) Sex limited character
 - (4) Autosomal dominant
- **Q.136**Which one of the following is associated with sex-linked inheritance:
 - (1) Night-blindness
- (2) Haemophilia
- (3) Astigmatism
- (4) Polydactyly
- Q.137 Haemophilic female marries normal male, the theoretical ratio of their offsprings regarding haemophilia will be:
 - (1) All offsprings are haemophilic
 - (2) All girls are haemophilic
 - (3) All sons are haemophilic
 - (4) Half daughters and half sons are haemophilic
- **Q.138**If a colourblind woman marries with a normal man. The offspring will be:
 - (1) All colourblind
 - (2) All daughters will be normal and all son will be colourblind
 - (3) All normal
 - (4) All daughters will be colourblind and all sons will be normal
- Q.139 If a colourblind man is married to a normal woman (Absence of gene of colourblindness on its X-chromosomes), then from the point of view of disease, their offsprings will be:
 - (1) All sons will be colourblind
 - (2) All daughters will be colourblind
 - (3) All sons and daughters will be normal
 - (4) All sons and daughters will be colour blind
- Q.140 If a colour blind man marries a girl who is normal (homozygous) for this character, then genotypically:
 - (1) Sons and daughters will be normal
 - (2) Sons will be colour blind, daughters will be normal
 - (3) Sons will be normal, daughters will be carriers
 - (4) Both sons and daughters will be colour blind
- **Q.141**Frequency of crossing over will be relatively more if:
 - (1) Distance between the two genes is less
 - (2) Distance between the two genes is more
 - (3) Linked genes are more

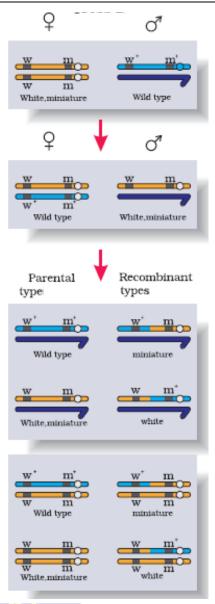


- (4) Both (2) and (3)
- Q.142 Presence of recombinants is due to:
 - (1) Crossing over
 - (2) Linkage
 - (3) Lack of independent assortment
 - (4) All of the above
- **Q.143**Morgan coined the term_____ to describe the physical association of genes on a chromosome and the term_____ to describe the generation of non-parental gene combinations :
 - (1) Recombination; Linkage
 - (2) Recombination; Non-recombination
 - (3) Linkage; Non-recombination
 - (4) Linkage; Recombination
- **Q.144**Experimental verification of the chromosomal theory of inheritance done on the basis of work with:
 - (1) Pea plant
- (2) Sweet pea plant

- (3) Snapdragon
- (4) Drosophila
- Q.145 Morgan and his group found that when genes were grouped on the same chromosome, some genes were very tightly linked and showed:
 - (1) Very low recombination
 - (2) Higher recombination
 - (3) No recombination
 - (4) 100% parental combination

Q.146





On the basis of above cross, the percentage of parental type in F_2 generation is

- (1) 98.7% (2) 37.2% (3) 62.8% (4) 1.3%
- **Q.147** The experimental verification of the chromosomal theory of inheritance was given by:
 - (1) Boveri
- (2) Sutton
- (3) T.H. Morgan
- (4) Bateson

SEX DETERMINATION

- **Q.148**Initial clue about the genetic/chromosomal mechanism of sex-determination can be traced back to some of the experiments carried out in:
 - (1) Human beings
- (2) Birds
- (3) Insects
- (4) Plants
- **Q.149**In which of the sex determination both male and female have same number of chromosomes?



(1) XY type (2) ZO type (1) ZW female in birds (3) XO type (4) Both (1) and (3) (2) XX female in *Drosophila* (3) XX female in Grasshopper Q.150 Two different types of gametes in terms of the sex (4) ZZ male in birds chromosomes are produced by: Q.158 Which of the following symbols are used for (1) Female fruit fly representing sex chromosome of birds: (2) Male butterfly (2) XX - XY(1) ZZ - ZW(3) Male human and female Drosophila (3) XO - XX(4) ZZ - WW(4) Female birds **Q.159**in honeybee males produce sperms by **O.151** Individuals having homomorphic sex (1) Meiosis (2) Mitosis chromosomes produce: (3) Amitosis (1) Only one gamete in complete life span (4) More than one option correct (2) One type of gametes **Q.160** Sex determination in humans takes place by: (3) No gametes (1) Sex chromosomes of father (4) Two type of gametes (2) Measurement of sperm (3) Measurement of ovum Q.152 How the sex of offsprings determined in humans: (4) Sex chromosomes of mother (1) By sex chromosome of mother Q.161 The number of chromosomes in female & male (2) By size of ovum respectively in honeybee is (3) By size of sperm (1) 16, 16(2) 16, 32(4) By sex chromosome of father (3) 32, 16 (4) 32, 32 Q.153 Which of the following possess homogametic Q.162 Which of the following is responsible for sex male: determination in chick: (1) Plants (2) Man (1) Sperm (2) Egg (3) Insect (4) Birds (4) Every cell of body (3) Somatic cell Q.154Which chromosome set is found in male grass **Q.163** In which of the following sex is determined by hopper: female individual: (1) 2A + XY(2) 2A + XO(1) Human (2) Drosophila (3) 2A + YY(4) 2A + XX(3) Birds (4) Grasshopper **Q.164** Male heterogamy found in case of: Q.155In *Drosophila*, the sex is determined by: (1) XO type male in Grasshopper (1) The ratio of number of X-chromosomes to (2) XY type male in human the sets of autosomes (3) ZW male in birds (2) X and Y chromosomes (4) 1 and 2 both (3) The ratio of pairs of X-chromosomes to the **Q.165** In which of the following monosomic male is pairs of autosomes found: (4) Whether the egg is fertilized or develops (1) Human (2) Birds parthenogenetically (4) Grasshopper (3) Honey bee Q.156In honey bee, which of the following isn't formed **GENETIC DISORDERS** by fertilization? Q.166In pedigree analysis, symbol given for sex (1) Queen unspecified is: (2) Workers (2) (3) Drones (4) More than one option is correct. Q.167In the given pedigree, Indicate whether the

shaded symbols indicate dominant or recessive

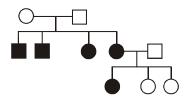
allele:



condition

Q.157 Which of the following represent heterogametic

PRINCIPLES OF INHERITANCE AND VARIATION



- (1) Recessive
- (2) Codominant
- (3) Dominant
- (4) It can be recessive or dominant both
- **Q.168**Match the following (w.r.t. Pedigree analysis):

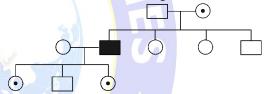
Column -I

Column -II

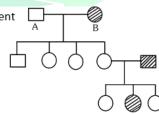
- a. Solid symbol
- (i) Carrier of sex linked trait
- b. Horizontal line between symbols
- (ii) Offspring
- (iii) Trait to be studied c. Horizontal line above the symbols
- d. Dot in centre
- (iv) Parents
- (1) a(iv), b(iii), c(ii), d(i)
- (2) a(ii), b(iii), c(iv), d(i)
- (3) a(iii), b(iv), c(ii), d(i)
- (4) a(i), b(ii), c(iv), d(iii)
- Q.169 The inheritance of a particular trait is represented in the family tree over generations in
 - (1) Pedigree analysis (2) Linkage
 - (3) Recombination (4) Mutation
- **Q.170** If a cross is made between two individuals each having genotype Bb, two offsprings are obtained. Out of these first has dominant trait. What is the probability that the second offspring will exhibit recessive trait:
 - (1) 1/4
- (2) 100
- (3) Zero (4) 3/4
- Q.171 Which one of the following character in man is controlled by recessive gene:
 - (1) Colourblindness
 - (2) Myotonic dystrophy
 - (3) Brachy-dactyly
 - (4) Curly hairs
- Q.172 This symbol represents
 - (1) Sex specified
 - (2) Five unaffected offsprings
 - (3) Mating
 - (4) Normal male

- Q.173 Failure of segregation of chromatid during cell division cycle results in the gain or loss of a chromosome(s), called
 - (1) Aneuploidy
 - (2) Polyploidy
 - (3) Euploidy
 - (4) More than one option is correct
- Q.174Down's syndrome is caused by an extra copy of chromosome number 21. What percentage of offsprings produced by an affected mother and a normal father would be affected by this disorder:
 - (1)50%
- (2) 25%
- (3) 100% (4) 75%
- **Q.175** A male human is heterozygous for autosomal genes A and B and is also hemizygous for haemophilic gene h. What proportion of his sperms will be abh:

- (1) $\frac{1}{4}$ (2) $\frac{1}{8}$ (3) $\frac{1}{32}$ (4) $\frac{1}{16}$
- **Q.176**Predict from the following chart:

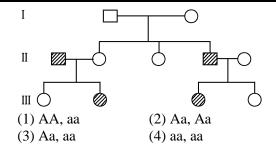


- (1) Character is dominant and carried by X chromosome
- (2) Character is carried by Y chromosome
- (3) Character is sex linked recessive
- (4) Character is autosomal recessive
- **Q.177** Given pedigree shows inheritance of autosomal recessive gene. What is the genotype of given parents:

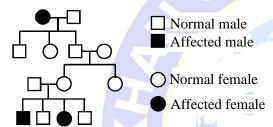


- (1) AA, aa
- (2) aa, AA
- (3) aa, Aa
- (4) Aa, Aa
- Q.178A pedigree is shown below for a disease that is autosomal recessive. The genetic make up of the first generation:

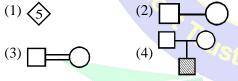




- **Q.179** Which of the following is incorrect about thalassemia?
 - (1) Autosomal recessive trait
 - (2) Quantitative inheritance
 - (3) Two types α & β thalassemia
 - (4) Shows pleiotropy
- Q.180 Study the pedigree given below and assign the type of inheritance of the trait:



- (1) X-linked recessive
- (2) Y-linked
- (3) Autosomal recessive
- (4) Autosomal dominant
- **Q.181** Which of the following symbol is used for mating between relatives (Consangeineous mating):



- **0.182** Cystic fibrosis, Haemophilia and Thalassemia are:
 - (1) Chromosomal disorders
 - (2) Autosomal recessive disorders
 - (3) Mendelian disorders
 - (4) Autosomal dominant disorders
- **Q.183**Phenylketonuria is an inborn error of metabolism that is inherited as:
 - (1) Autosomal recessive trait
 - (2) Sex-linked dominant trait
 - (3) X-linked recessive trait
 - (4) Autosomal dominant trait
- **Q.184**Which of the following abnormalities is due to autosomal dominant mutation?

- (1) Colour blindness
- (2) Thalassemia
- (3) Myotonic dystrophy
- (4) Haemophilia
- **Q.185** Absence or excess or abnormal arrangement of one or more chromosomes results in :
 - (1) Point mutation
 - (2) Chromosomal disorders
 - (3) Mendelian disorders
 - (4) Gene mutation
- Q.186Mark the odd one w.r.t. syndrome which occur due to failure of segregation of homologous pair chromosomes during cell division cycle.
 - (1) Klinefelter's syndrome
 - (2) Down's syndrome
 - (3) Turner's syndrome
 - (4) Thalassemia
- **0.187** Mark the correct match:
 - (1) Turner's syndrome 46 + XO
 - (2) Phenylketonuria 44 + XYY
 - (3) Klinefelter's syndrome 44 + XXY
 - (4) Thalassemia 44 + YO
- Q.188In which of the following disorders affected individuals possess 47 chromosomes?
 - (1) Turner's syndrome
 - (2) Klinefelter's syndrome
 - (3) Down's syndrome
 - (4) Both (2) and (3)
- **Q.189**In which of the following disorder gynecomastia symptom is seen in individuals?
 - (1) Down's syndrome
 - (2) Turner's syndrome
 - (3) Klinefelter's syndrome
 - (4) Phenylketonuria
- Q.190 Allosomic trisomy condition is seen in:
 - (1) Turner's syndrome
 - (2) Klinefelter's syndrome
 - (3) Down's syndrome
 - (4) Both (2) and (3)

>>>>>>>



ANSWER KEY

TOPIC WISE QUESTIONS

Que.	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Ans.	2	4	3	1	4	2	2	2	4	2	4	2	1	4	3
Que.	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30
Ans.	3	3	3	4	4	4	2	1	4	4	3	3	1	4	1
Que.	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45
Ans.	1	1	2	4	4	3	1	1	2	1	2	2	1	2	3
Que.	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60
Ans.	3	1	1	3	3	2	1	3	2	1	1	2	2	4	2
Que.	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75
Ans.	1	3	3	1	4	4	3	3	2	4	2	3	4	4	1
Que.	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90
Ans.	2	3	2	3	2	2	3	3	4	4	3	2	2	4	3
Que.	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105
Ans.	1	3	1	3	4	3	2	4	3	3	2	4	3	1	1
Que.	106	107	108	109	110	111	112	113	114	115	116	117	118	119	120
Ans.	1	1	2	4	4	1	3	1	1	3	1	2	4	3	2
Que.	121	122	123	124	125	126	127	128	129	130	131	132	133	134	135
Ans.	1	2	2	2	3	2	3	1	1	2	2	3	2	4	1
Que.	136	137	138	139	140	141	142	143	144	145	146	147	148	149	150
Ans.	2	3	2	3	3	2	1	4	4	1	3	3	3	1	4
Que.	151	152	153	154	155	156	157	158	159	160	161	162	163	164	165
Ans.	2	4	4	2	1	3	1	1	2	1	3	2	3	4	4
Que.	166	167	168	169	170	171	172	173	174	175	176	177	178	179	180
Ans.	1	1	3	1	1	1	2	1	1	2	3	1	2	4	3
Que.	181	182	183	184	185	186	187	188	189	190					
Ans.	3	3	1	3	2	4	3	4	3	2					

Usted Learning

** KHAN SIR **

