



Tips & Tricks Questions (2025-2026)

Class-XII

Subject: Biology

Chapter Name : Principles of Inheritance and Variation (Chapter-04)

1. Master Mendel's Crosses (Very High Weight-age)

- Learn one base cross (monohybrid & dihybrid) and derive others from it.
- Always write:
 - P generation
 - Gametes
 - Punnett square
 - Genotypic & phenotypic ratio
- Trick:
 - Monohybrid phenotypic ratio $\rightarrow 3 : 1$
 - Dihybrid phenotypic ratio $\rightarrow 9 : 3 : 3 : 1$CBSE checks steps, not just the final ratio, don't use short forms

2. Understand, Don't Simply Memorize Laws of Inheritance

- Law of Dominance \rightarrow Monohybrid cross
 - Law of Segregation \rightarrow Purity of gametes
 - Law of Independent Assortment \rightarrow Dihybrid cross
- Link law
- \rightarrow
- experiment
- \rightarrow
- ratio

3. Pedigree Analysis

- Square = male, Circle = female
 - Shaded = affected
 - Autosomal dominant \rightarrow appears in every generation
 - Autosomal recessive \rightarrow may skip generations
 - X-linked recessive \rightarrow more males affected
- Trick : "Only males suffer and females are carriers
- \rightarrow
- X-linked"

4. Linkage & Recombination

- More distance between genes \rightarrow more recombination
 - Linked genes \neq independent assortment
- Remember : Morgan's experiments proved linkage, not Mendel

5. Practice Numerical Problems

- Blood group inheritance
 - Sex-linked inheritance (haemophilia, colour blindness)
 - Test cross vs back cross
- Remember : Test cross ALWAYS involves homozygous recessive

6. NCERT Diagrams Are Mandatory

- Meiosis (segregation of alleles)
- Pedigree charts
- Chromosomal theory of inheritance

MENDELIAN DISORDERS

- Due to mutation in a single gene
 - Autosomal Recessive \rightarrow "PAST"
Phenylketonuria, Albinism, Sickle cell anaemia, Thalassemia
- Trick: These disorders often "hide" and appear when both parents are carriers.

- Autosomal Dominant → “**H**”
Huntington’s disease
Trick: Dominant = appears in every generation
- X-Linked Recessive → “**He Cares**”
Haemophilia, Colour blindness
Trick: More males are affected, females are usually carriers

Gene Location Trick

- X-linked → sex chromosomes
- Autosomal → chromosome 1-22
Trick: If males suffer more → X-linked

CHROMOSOMAL DISORDERS

- Due to abnormal number or structure of chromosomes

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- Down’s syndrome
- Turner’s syndrome
- Klinefelter’s syndrome

Remember by Chromosome Formula

Disorder	Chromosome Number	Memory Trick
Down’s syndrome	Trisomy 21	
Turner’s syndrome	XO (45)	Turner = Only One X
Klinefelter’s	XXY (47)	Extra X in male

- Turner’s → only females; Klinefelter’s → only males
- Down’s syndrome → nondisjunction
- Turner’s / Klinefelter’s → sex chromosome nondisjunction

COMMON ERRORS

- × Calling Down’s Syndrome a Mendelian disorder
- × Writing haemophilia as autosomal
- × Forgetting chromosome number (45/47/21)
- × Saying dominant = more common in population
- × Writing Ratio Without Showing Cross
- × Confusing Genotype & Phenotype
- × Mixing Up Mendel & Morgan

Mendel	Morgan
Pea plant	Drosophila
Independent assortment	Linkage
Phenotypic ratios	Gene mapping