

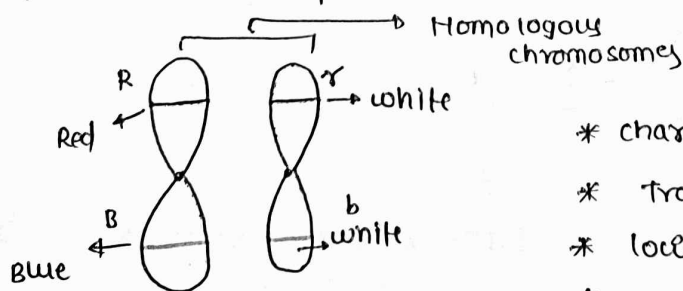
17/1/20

GENETICS - 1

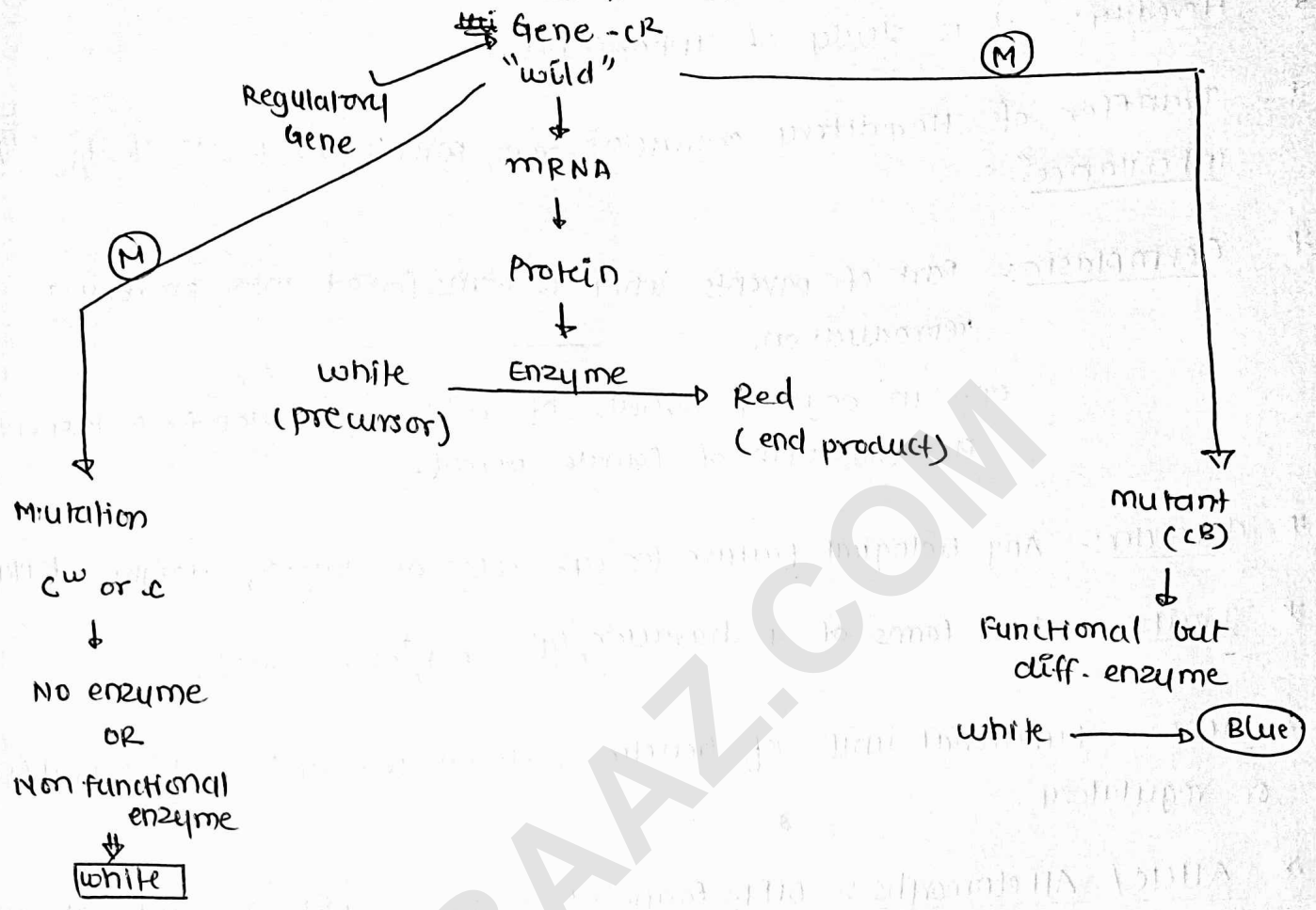
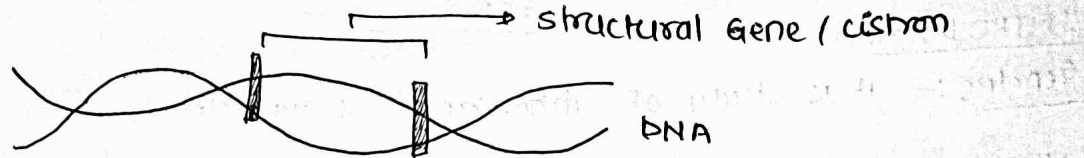
(I) INTRODUCTION

- * Genetics :- It is study of inheritance & variations.
- * Heredity :- it is study of inheritance.
- * Transfer of Hereditary characters from parents to progeny is klg inheritance.
- * Germplasm :- Part of parents which is transferred into progeny in sexual reproduction.
eq:- in oogamy nucleus of σ^2 + ϕ Gametes + Mitochondrial & Plastidial DNA of female parent.
- * Character :- Any biological feature for eq:- color of flower, Height of plant etc.
- * Traits :- diff. forms of a character, eq:- Red, Blue, white;
- * Gene :- Functional unit of genetic material (it may be structural (eiston), or regulatory).
- * Allele / Allelomorphs :- diff forms of a gene which are int at same locus but produce diff traits
- * A Gene may have many Alleles.
- * Gene control characters by producing proteins. A Gene may control many characters & a character may be controlled by ~~may~~ many genes.
- * Locus :- Address or position of a gene.

eq:- In foll. diagram there are



- * characters = 1, colour of flower
- * traits = Blue, Red; white
- * loci/genes = 2 (R & B)
- * Alleles \rightarrow R has 2 Allele \rightarrow R, r
B " " " \rightarrow B, b



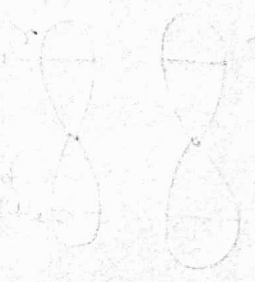
* Genotype $\xrightarrow{+ \text{environment}}$ Phenotype
 eq: - TT \rightarrow eq: - Tall

* Possible Genotypes = $\frac{n(n+1)}{2}$ for 2n

n = Types of Alleles.

* Abo system has 3 Alleles.
 (in humans)

so possible Genotypes = $\frac{3 \times 4}{2} = 6.$



Types of Gametes

Q. How many types of Gametes can be produced from $AaBbCc$

Ans.

$$\text{Type of Gametes} = 2^n$$

$n = \text{no. of Hybrid}$

$$AaBbCc = 2^3 = 8$$

$$A-B \begin{cases} C \\ c \end{cases} = \begin{matrix} ABC \\ ABc \end{matrix}$$

$$a-b \begin{cases} C \\ c \end{cases} = \begin{matrix} aBC \\ abC \end{matrix}$$

① $RrTt$
 $2^2 = 4$

② $aabb = 2^1 = 2$

③ $aabb = 2^0 = 1$

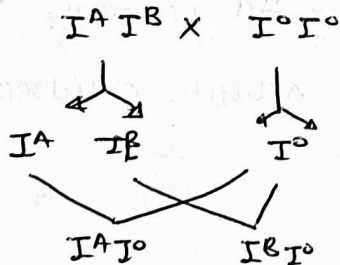
④ ~~Genotype of Mohan~~
is $AaBb$
due to human (O^+)
 $AaBb \times Y = 2^3 = 8$

* Use of Probability ; Permutation & combination

in Qs.

"AND" \longrightarrow "X" Multiply the probabilities
"OR" \longrightarrow "+" Addition

Ques:- Blood group of parents is $I^A I^B$ & $I^O I^O$
Probability of boy with B Blood group.



Probability of σ^+ = $\frac{1}{2}$
" " B D. = $\frac{1}{2}$
so σ^+ of B = $\frac{1}{4}$

Q. A couple has a child what is probability of its being boy or girl

Ans. $\frac{1}{2} + \frac{1}{2} = 1$

TERY X TERY

P. of Tall + white.
T d R w
(3:1) (3:1)

20/10/16

L-2

TL-79

Q. A couple has 4 children, what is possibility of one boy + 3 girls?

Sol.

$$\frac{n!}{n_1! \times n_2!} (P_1)^{n_1} (P_2)^{n_2}$$

n = Total events = 4

n₁ = 1 (Boy)

P₁ = Probability of boy = $\frac{1}{2}$

n₂ = 3 (Girl)

P₂ = Probability of Girl = $\frac{1}{2}$

$$= \frac{4!}{1! \times 3!} \left(\frac{1}{2}\right)^1 \left(\frac{1}{2}\right)^3$$

$$= \frac{4 \times 3 \times 2 \times 1}{1 \times 3 \times 2 \times 1} \times \frac{1}{2} \times \frac{1}{8} = \frac{1}{4}$$

Q. 44

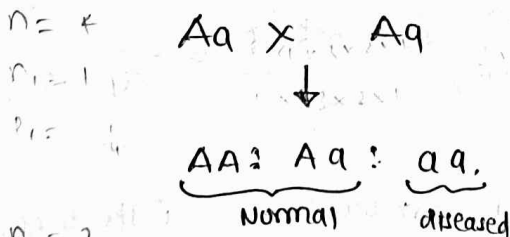
Albinism in human is controlled by "a" (recessive gene), both (aa)

• parents are carriers / heterozygous - Aa (Normal)

what is possibility of 1 Normal + 3 Albinic children in family of

4.

Sol.



n₂ = 3
n = 4

n₁ = 1 (Normal)

n₂ = 3 (Albinic)

P₁ = $\frac{3}{4}$

P₂ = $\frac{1}{4}$

$$= \frac{4!}{1! \times 3!} \times \left(\frac{3}{4}\right) \left(\frac{1}{4}\right)^3 = \frac{3}{64}$$

* Genotypic ratio \rightarrow $RR : Rr : rr$
 $\quad\quad\quad 1 : 2 : 1$
 Phenotypic Ratio Red : white
 (= Monohybrid Ratio) $3 : 1$

Note :- * Parents =? from the starting of cross

* Phenotype is preferred over Genotype if not mentioned in the Q.

* $RR \times rr \rightarrow Rr$
 $Rr \times Rr \rightarrow RR : Rr : rr (1:2:1)$
 $Rr \times rr \rightarrow Rr : rr (1:1)$
 $Rr \times RR \rightarrow Rr : RR (1:1)$

(B) Dihybrid cross :- study of any two character

eg:- Height of plant + colour of the flower.

P_1 :- Tall, Red \times Dwarf white

G_1 :- $TTRR$ \times $ttrr$

F_1 :- TR \times tr

$TtRr$ Tall, red

P_2 : $\sigma^{\text{♂}}$ $TtRr$ Tall red \times ♀ Tall red $TtRr$

* Results of F_2 Generation

Genotype	G. Ratio	Phenotype	Ph. Ratio
$TTRR$	$2^0 = 1$	Tall Red $T_R_$	9 Parental
$TTRr$	$2^1 = 2$		
$TtRR$	$= 2$		
$TtRr$	$= 4$		
$TTrr$	$= 1$	Tall white T_rr	3
$Ttrr$	$= 2$		

tt RR	= 1	Dwarf Red	3
tt Rr	= 2	ttR_	
tt rr	= 1	Dwarf white	1 (Parental)
		tt rr	

In F₂ Gen. (n=2 ; dihybrid cross)

- Total phenotypes = $(2)^n = 2^2 = 4$
- Phenotypic Ratio = $(3:1)(3:1) = 9:3:3:1$
- No. of boxes in checker board = $(4)(4) = 16$
(Total zygote)
- Genotype = $(3)(3) = 9$
- Genotypic Ratio = $(1:2:1)(1:2:1)$
=

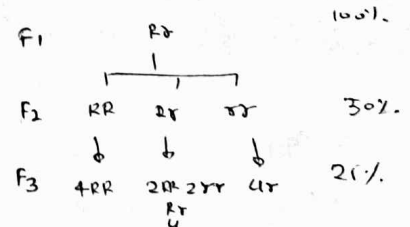
★ Tri hybrid cross (in F₂ Gen.)

- $2^3 = 8$
- 27 : 9 : 9 : 3 : 9 : 3 : 3 : 1
- $(4)^3 = 64$
- Genotype → $3 \times 3 \times 3 = 27$

Q. what is the % of hybrid in F₂ Gen. of a monohybrid cross?

Sol. % of Hybrid in

F₁ → F₂ → F₃ → F₄
100% 50% 25% 12.5%



Q. In a typical dihybrid cross 800 plants are produced what is the no. of Tt Rr plant.

$$\frac{4}{16} \times 800 = 200$$

$$\frac{2}{4} \times \frac{2}{4}$$

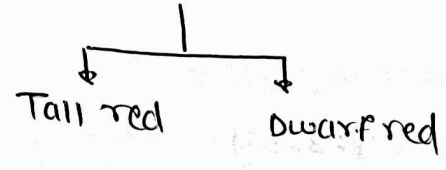
* Parental : Recomb = 10:16

- Pure — 4/16
- Monohybrid — 8/16
- Dihybrid — 4/16

Q. A Tall plant with Red flowers is crossed with dwarf plant with white flowers + produce 100 Tall red + so dwarf ~~white~~ red
 what is Genotype of parents

Sol.

$$T_R_ \times tt\ rr$$



$$TtRr$$

Q. A Tall plant with Red ^{dominant} Axillary flowers is crossed with dwarf plant having white + terminal flower + it produce all Tall plant with red flowers 50% plants are terminal + 50% with Axillary flower

$$TTR_ A_ \times tt\ rraa$$

$$T_ R_ aa$$

$$TTRRAa$$

$$TTRRAa \times tt\ rraa$$

$$T_ R_ aa$$

$$T_ R_ Aa$$

Q. A Person with AB B.G marries a women of B B.G whose father was O what is possibility of B.G in their child

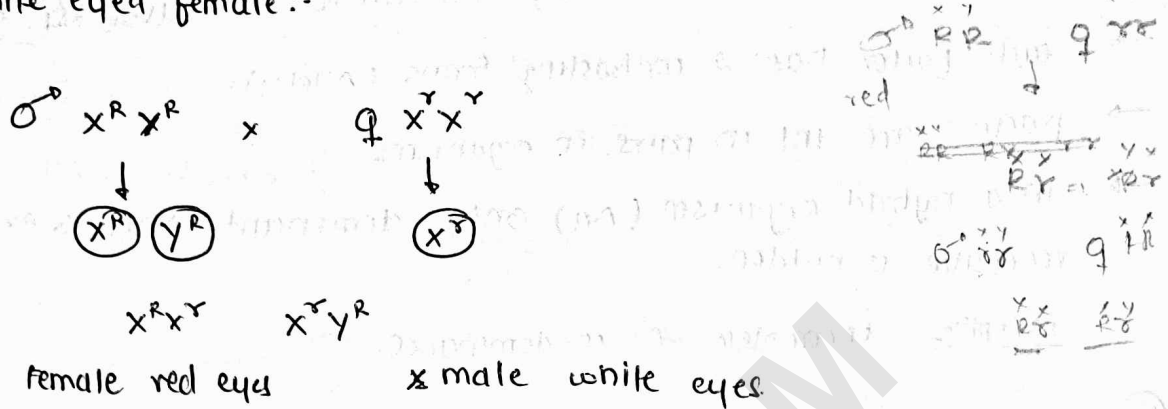
$$AB \times BO$$



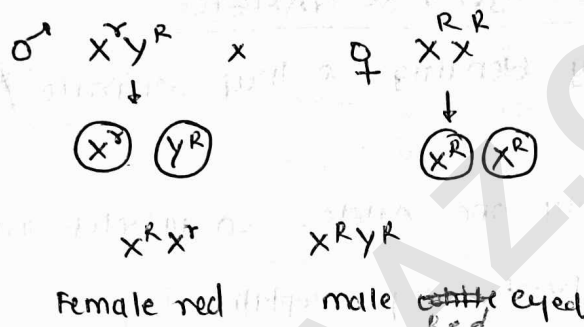
① Reciprocal crosses

Q. in *Drosophila* color of eyes is x linked + red color is dominant over white, what is phenotype of offspring if red eyed male is crossed with white eyed female?

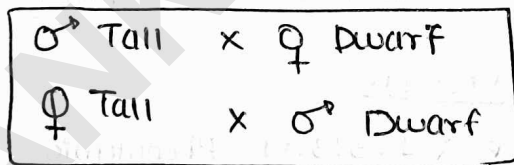
Sol.



Sol.



* in R.C parents i.e source of gametes are inter changed to observe the effect of sex of parent on the phenotype of offsprings.
for eg:-



* results are changed in cytoplasmic + sex linked inheritance but results remain unchanged in mendelian inheritance (i.e Nuclear + Autosome).

① Laws of Mendel

- on basis of hybridisation experiment in garden pea (*Pisum sativum*; $2n=14$)

① LAW OF DOMINANCE

- characters are controlled by particulate structures k/a "factors" (Gene)
- each factor has 2 contrasting forms (Alleles).
- factors are not in pairs in organisms.
- in a hybrid organism (Aa) only dominant trait is expressed & recessive is hidden.
- exceptⁿ:- incomplete & co-dominance.

② LAW OF SEGREGATION / PURITY OF GAMETES

- factors do not show any blending & they separate / segregate during gametogenesis
- each gamete receives only one allele. so gametes are always pure
- universal law i.e. without any exception.
- * segregatⁿ occurs during Meiosis. (Anaphase-1) so during gametogenesis in animals & spermatogenesis in plants.
- * Above mentioned laws were based on monohybrid cross i.e. 3:1 ratio.

③ LAW OF INDEPENDENT ASSORTMENT

- based on dihybrid cross i.e. (9:3:3:1) Phenotypic ratio
- Acc. to this law inheritance of 1 character remains independent of other characters.
- exception:- linkage
- explanation:- gene not on same chromosome (Homologous) show linkage i.e. inherit together while gene not on diff chromosome (Heterologous) show independent assortment.

Q. Opp. phenomenon of linkage is

- (a) independent Assortment
- (b) crossing over

Q. Occurrence of recessive phenotype in F_2 generation is explained by

- (a) law of segregation
- (b) law of purity of Gametes

24/1/20

L-4

TL-81

④ TEST AND BACK CROSS

Back cross

F_1 Hybrid

eg:- $AaBb$

x

$AA BB$ or $aa bb$.

(Parental genotype in typical cross)

Test cross

unknown Genotype x Recessive

OR

$Aa \times aa$

Backcross / outcross

eg:- $Aa \times AA$ (Pure dominant)

→ To suppress the recessive Allele

OR

To test Potential Genotype

Significance:-

→ To determine the Genotype

OR

linkage.

* Each test cross is a back cross but back cross is not a test cross.

* Test cross ratio (Typical)

- Monohybrid ($Aa \times aa$) = 1:1
- Dihybrid = 1:1:1:1
- Trihybrid = 1:1:1:1:1:1:1:1

Q. No. of characters in Pea Pod → 3 [Pod shaped, color, flower position]

Q Back cross is

1) $AA \times aa$

2) $Aa \times Aa$

3) $AA \times aa$ (Test cross)

4) $Aa \times AA$

} BACK cross.

5) GENE INTERACTIONS

A) Intra-Genic / Inter Allelic

blw Allele of a Gene
(same cross)

eg:- incomplete dominance



A) Intra-Genic / inter Allelic

① Incomplete Dominance

* Discovered by Correns in *Mirabilis*

* In this phenomenon dominant gene can't suppress phenotype of recessive Allele so an intermediate phenotype in F_1 generation.

* F_2 Generation is not similar to any parent.

* Test cross is not required.

* eg:- pink color flower in

- *Mirabilis Jalapa* (4 o'clock plant)
- *Antirrhinum majus* (snapdragon / dog flower)

Red Flower x white
(RR) (rr)



F_1 Rr (PINK)



RR Rr rr

1 : 2 : 1

Red

Pink

ratio.

→ same Genotype + same phenotypic ratio.

Q. A pure red flowered plant is crossed with a white flowered plant. what is % of red flowered plant in F_2 generation.

- (A) 25 (B) 50 (C) 75 (D) 100

Q. IN a ~~series~~ species, character of Height show complete dominance (tall & dwarf) and character of Flower color show I.C. Dominance (Red, Pink & white) 800 plants are produced in F_2 generation then:

- (A) F_2 phenotypic ratio
(B) No. of tall, pink plants
(C) No. of white flowered plants
(D) No. of TtRr plants.

Sol. (A) $\begin{matrix} T & P & R & P & W \\ (3:1) & (1:2:1) \\ 3:6:3 & : & 1:2:1 \end{matrix}$

(B) $\frac{6}{16}$ or $\frac{3}{4} \times \frac{2}{4} = \frac{6}{16} \times 800 = 300$

(C) $\frac{4}{16} \times 800 = 200$

(D) $\frac{2}{16} \times 800 = 100$

Ques. what is ratio of any one pair of contrasting character in dihybrid cross?

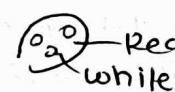
Sol:- 3:1

Ques:- what is ratio of 2 pair of contrasting characters in a trihybrid cross?

Sol. 4:3:3:1

② codominance (1:2:1)

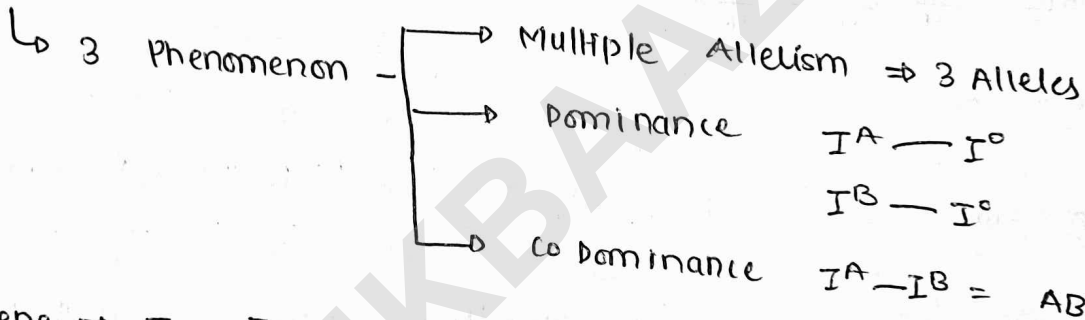
↳ Both Alleles express themselves equally so F₁ show mosaic of both phenotypes

- eg:-
- SKIN color in CATTLE (Red x white → Roan  } Resemblance to both parents
 - AB blood grp ; examples of multiple Allelism.

Q. same Genotypic + Phenotypic ratio are found in:-

- a) incomplete } both (if given)
- b) codominance }
- c) dominance
- d) complementary gene

★ ABO Blood Group system



Gene → I = Iso agglutinogens

OR

L = Discovered by Landsteiner + Levine.

[L^A + L^B or L^o/l]

→ on 9th chromosome

• Rh factor on 1st chromosome

Ques:- ABO B-G syst. is controlled by three Alleles what is possible no. of Alleles in population, somatic cells + Gametes?

Sol.



Ques. what is no. of possible genotypes in ABO B-G-s in humans.

Sol.

Acc. to 2N

$$= \frac{3(3+1)}{2} \Rightarrow 6$$

③ Multiple Allelism

occurrence of more than 2 Alleles for eq.

- sexual incompatibility
- ABO Blood Group syst.
- Coat color in most of animals (eq: cow, dog, Rabbit)
- eye color in *Drosophila*.

④ Pleiotropism

A Gene affects two or more characters

eq:- Most of genetic disorders, lethal Genes + 'B' Gene of Garden Pea.

* ~~B = Branching~~

* B = starch branching enzyme

Genotype	starch Granules	shape of seed
BB	Larger	Round
Bb	Medium (incomplete domin.)	Round (Dominance)
bb	smaller	wrinkled

starch Granules → incomplete dominance

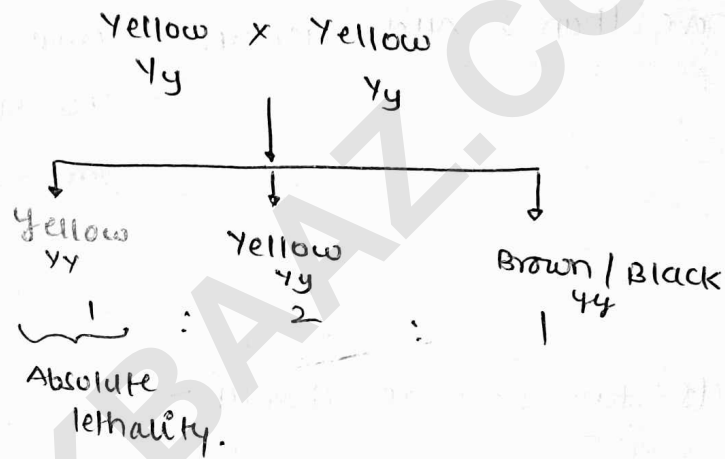
shape of seed → complete dominance.

⑥ lethal Gene

- * These genes cause death or severe disability in organism
- * lethality may appear
 - before birth (Absolute lethality)
 - In childhood (sub lethality)
 - After sexuality mature (delayed lethality)
- * lethal Gene are recessive in nature & also show pleiotropism.

* foreq:- • many genetic disorders

• Yellow coat color in mice



* "y" gene is dominant for skin color
 recessive for lethality.

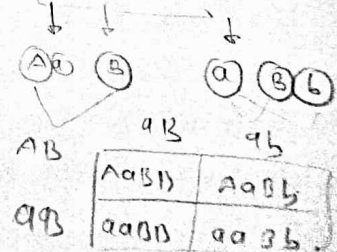
Q. } Flower color in a plant is controlled by 2 genes. Presence of any dominant Allele produce red color
 white color appears when both genes are in recessive form.
 presence of both gene also produce red color.

① what is phenotypic cross in ~~parent~~ F₂ generation

" " " " if parents are

A_ B_
 aa B_

AaBB \times aaBb.



Hint:- Duplicate Gene

Q. In *Lathyrus odoratus* (sweet pea) flower color is controlled by 2 genes

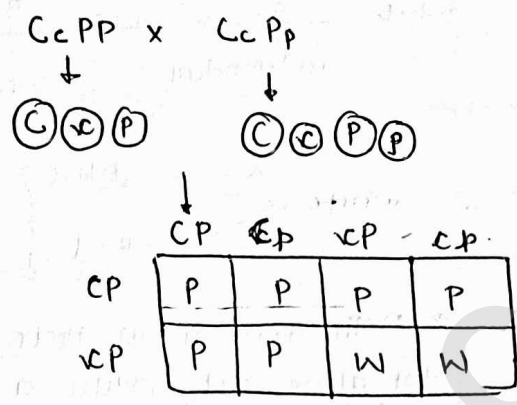
- Purple color appears if both genes are present
- in other combination color remains white.

Q. (a) what is dihybrid ratio

(b) Phenotypic ratio if parents are $CcPp \times CcPp$.

(a) 9:7
Purple white

(b)



P.R = 9:7
P:W

दो Group हैं (C) & (P)
जब Pair होते तब
(C) - (P) -
blank में अगर कुछ भी रहेगा it will be dominant!
C - p -
अब Always white.

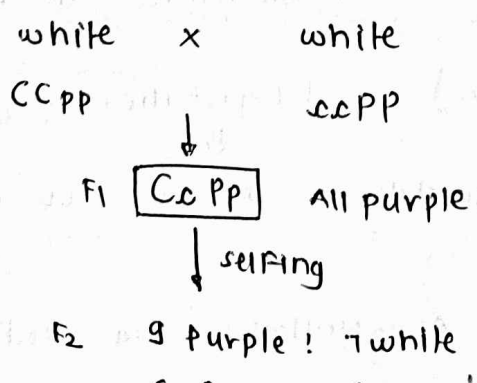
(B) Non-Allelic (or) Intergenic Interactions

these interaction occur b/w diff. gene i.e. Gene present at diff. loci

(1) Complementary Genes (9:7)

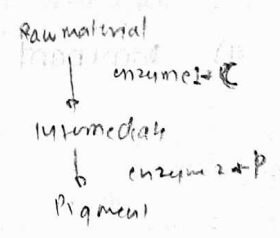
* Discovered by Bateson & Punnett while studying flower color in sweet pea.

* These genes show no phenotype or similar phenotype when present alone but interact to produce a new phenotype when present together.



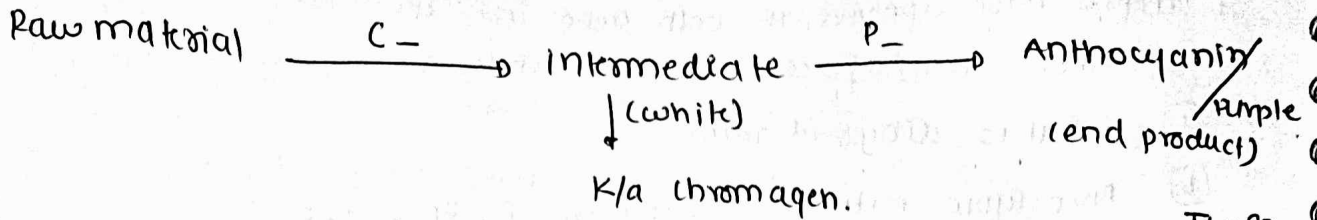
C - P - → Dominant
अब भी recessive आता → white

(3 C - Pp)
(3 cc P -)
(1 ccpp)



Dominant form (अब भी)
है to produce
particular character

explanation



29/01

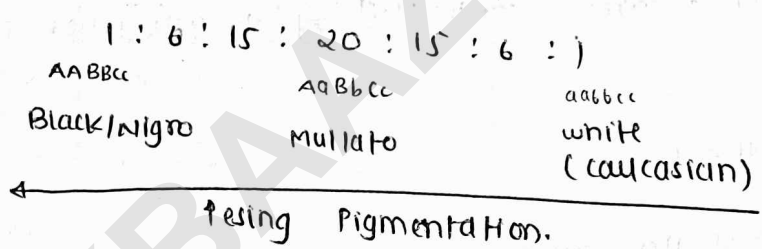
L-6

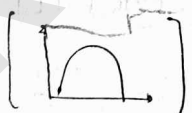
TL-82

Non Allelic G.I	Explanation
2) supplementary Genes (9:3:4)	white $\xrightarrow{A-}$ Blue $\xrightarrow{B-}$ Red independent dependent / Modifier eq: - coat color in mice
3) collaborative supp. Genes (9:3:3:1)	white $\begin{cases} \xrightarrow{A-} \text{Blue} \\ \xrightarrow{B-} \text{Red} \end{cases}$ } $A-B-$ Purple * Both Gene show their own phenotypes when tht alone but produce a new phenotype when tht together.
4) Duplicative Gene OR Pseudo Allely (15:1)	$A- = \text{Red}$ $aa = \text{white}$ $B- = \text{Red}$ $bb = \text{white}$ $A-B- = \text{Red}$ * similar behaviour at both loci.
5) Additive (OR) collaborative Gene (9:6:1)	* Duplicative Gene with additive effect $(A-B- = \text{Dark red})$
6) Inhibitory Gene (13:3)	white $I- \rightarrow x \begin{cases} \downarrow R- \\ \downarrow \text{Red} \end{cases}$ * $I-$ has no phenotype of its own but inhibits the expression of another gene.
7) Epistasis	similar to dominance but a <u>diff. loci</u> .
a) dominant epistasis (12:3:1)	$A- = \text{Red (epistatic)}$; $aa = \text{Neutral}$ Hypostatic $B- = \text{Blue}$; $bb = \text{white}$
b) Recessive Epistasis (9:3:4) * Ratio is similar to supplementary genes eq: - coat color in mouse 9:3:4 Agouti Black Albino.	$A- = \text{Neutral}$; $aa = \text{Red (epistatic)}$ Hypostatic. $B- = \text{Blue}$; $bb = \text{white}$

⑥ Quantitative (OR) Polygenic inheritance

- * discovered by Kolreuter
father of quantitative inheritance.
- * detailed study by Nilsson - Ehle in kernel/seed color.
- * controlled by 2 polygenic genes & ratio 1 : 4 : 6 : 4 : 1
(phenotypic) Red dark pink pink light pink white
- * each functional allele is k/a contributing allele & each ~~gene~~ is k/a "Poly Gene"
- * Polygene are duplicate gene.
- * Monohybrid ratio = 1:2:1
- * Dihybrid ratio = 1:4:6:4:1
- * Trihybrid ratio = for eg. skin color in human beings



- * environmental effect is more on quantitative character, so phenotype show cont. variations.  usually can't be characterised into discrete classes.

Que: - A Mullato (AaBbCc) marries with another Mullato what is % of Parental phenotype in offspring

→ $\frac{20}{64}$

Que: - Nigro marries with "caucasian" what is % of parental phenotype in their children?

6/ all mullatos

Que: in plant ht. is controlled by 2 polygenes. ht. of pure recessive (aabb) is 3 feet, ht. of pure dominant (AABB) is 11 ft & ht. of F₁ hybrid (AaBb) is 7 feet

(a) what is contribution of each polygene? (contribution Allele) - 2

(b) what is phenotypic ratio if both parents are AaBb

(c) what is PR if both parents are AaBb x AaBb.
1:3:3:1

Q. study of polygenic inheritance is more easier in plants due to
i) large no. offspring
ii) controlled + easing Matng.

* In human beings quantitative characters are studied in population!

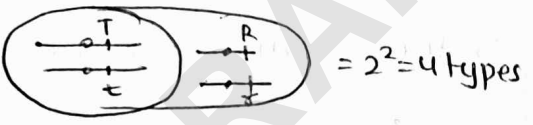
* In Humans color of eyes is controlled by 4 polygenes & height by 4 or more Polygenes.

7 BASICS OF LINKAGE

(A) Type of Pollen grain

If Genotype of PMC (TtRr)

Independent Assortment



(TR) : (Tr) : (tR) : (tr)
1 : 1 : 1 : 1

Linkage

complete linkage

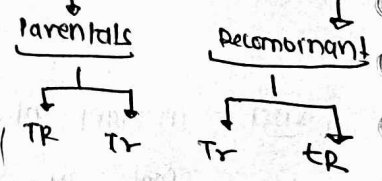
→ Gene exchange/co. =nt
→ No recombination in Test cross

(TR/tr) → TR : tr
2 : 2

(Tr/tR) → Tr : tR
2 : 2

Incomplete linkage

→ GE/co tnt
so Pollen from TR/tr PMC



• Gametes from Hybrid = Always 2 types
• " " Pure = Always 1 type

* In IC linkage % of Parentals + recombination varies but specific for particular characters of a species

Recombinant \leq Parental

$P_1 R_1 R_2 P_2$
3 : 1 : 1 : 3.

* Genotype of F₁ Hybrid

	P	R	RF	Distance	Parents of F ₁
AB/ab	AB:ab	Ab:aB	30%	30%	
+/--	+:--	-:-+	10%	10%	
AB/--	AB!--	A--B	20%	20%	
a+/+b	a+/+b	ab:++	10%	10mu	
+/ab	+:ab 40 40	+b:a+ 10 10	20%	20	
d ⁺ w ⁺ /dw	d ⁺ w ⁺ :dw 80 80	d ⁺ w:dwt 20 20	20%	20	

Q. In sweet pea, color of flowers + shape of pollen grains are controlled by two linked gene, from test cross of F₁ hybrid foll. plants were obtained.

Blue long = 10

Red round = 10

Blue round = 70

Red long = 70

Blue + long traits are dominant.

i) what is recombination freq.

ii) Distance

iii) crossing over freq.

iv) Genotype of F₁

v) Genotype of parents

$$\text{Recombination freq.} = \text{Distance} = \frac{1}{2}(\text{C.O.F})$$

C.O.F = crossing over freq.

sol. i) $\frac{20}{160} \times 100 = 12.5\%$

ii) 12.5%

iii) 25%

iv) B_L/bL

v) B_L/B_L + bL/bL

Ques. From a test cross of F₁ hybrid (d₁) tall white + Dwarf red plants are obtained in 1:1 ratio
 what is genotype of F₁ hybrid?

sol.

$$TtRr \times ttRR$$

$$\underline{TtRr} / \times \underline{ttrR}$$

T_r / t_r Tall white

t_r / t_r ~~Tall white~~ Dwarf red

Q. in sweet pea Homozygous blue flower (BB), long pollen plant (LL) go through cross with red flower round pollen (bb) (ll) (pure recessive) F₁ plant is test crossed + gives rise to 720 seed. How many of the offspring will be blue flower round pollen. Both gene are located at dist. of 12.5 mu.

sol.

$$\text{Dist.} = 12.5 \text{ mu} = \text{Recombination freq.}$$

$$12.5 = \frac{x}{720} \times 100$$

$$x = 90$$

$$\boxed{x = 45 \text{ plants}}$$

Trick

Monohybrid $1:2:1$

$1:3:3:1$

Dihybrid $1:4:6:4:1$

$1:5:10:10:5:1$

Trihybrid $1:6:15:10:15:6:1$

(B) CHROMOSOMAL BASIS OF INHERITANCE

In this section mechanism of inheritance is discovered and explained on basis of chromosome.

(A) chromosomal theory of inheritance (1902)

by Sutton + Boveri

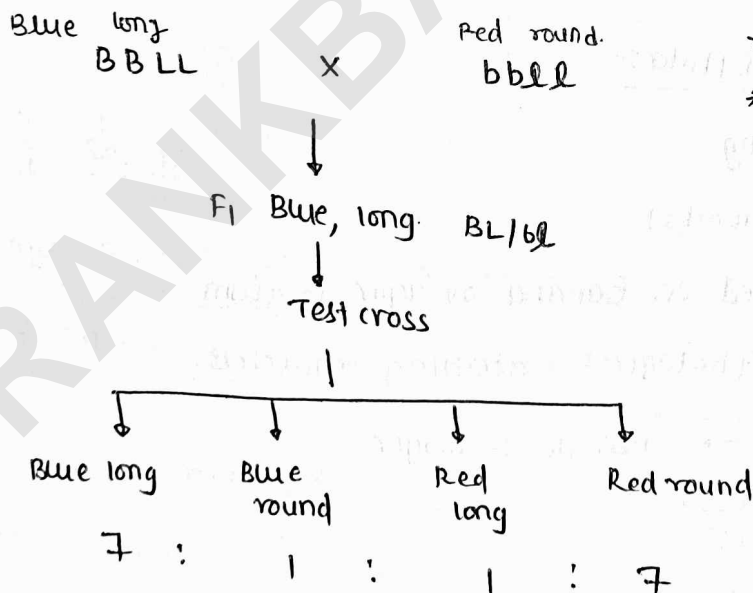
- "Mendelian factors (Genes), sit on chromosomes"
- it was based on 11th behaviour b/w Mendelian factors + chromosomes.
- it provide physical base to inheritance.
- chromosome are hereditary vehicle + connecting link b/w various generation.
- * "Inheritance of one pair is independent of other pairs" → Applicable to all chromosomes.

(B) ~~Gen~~

but not applicable to linked genes.

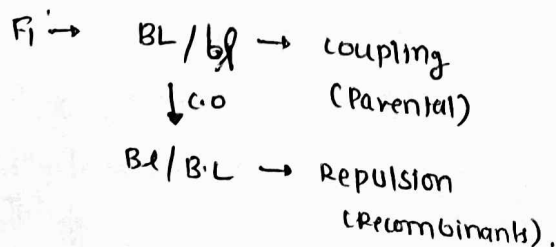
(B) coupling + repulsion hypothesis (1906-07)

- By Bateson + Punnett
- colour of flower and shape/size of pollen grains.



- * 1st linkage example
- * Autosomal + in plants.

Acc. to Morgan



① Chromosomal Theory of Linkage (1911)

* By T.H. Morgan et al

* Position of a gene is const. + specific + kly "locus".

* Gene's are arranged in a linear fashion

* Gene's tht on same chromosome inherit together

* linkage may be complete or incomplete.

* $\text{Recombination freq.} \propto \text{Distance}$

$\text{Strength of linkage} \propto \frac{1}{\text{Distance}}$
(% of Parental)

* TH Morgan is father of experimental genetics

* credit of discovery of linkage + crossing over

* study on Drosophila (fruit fly) ($2N=8$)

* Reason to choose Drosophila:-

- large no. of offsprings.
- life cycle short. (2 weeks)
- can be easily cultured on banana or agar medium
- presence of many morphological contrasting characters.
- sexual dimorphism → Female is larger

* Linkage Maps.

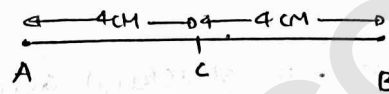
- * Based on recombination freq. of Test cross.
- so also k/a crossing over maps, genetic maps on chromosome maps.
- * 1st map was prepared by A. Sturtevant in drosophila. he was a student of Morgan
- * $1\% \text{ RF} = 1 \text{ mu} / 1 \text{ cM}$

Ques. What is genetic map of ABC Gene if R.F of various test cross are as follows:-

$$A \text{ \& } B = 10\%$$

$$B \text{ \& } C = 4\%$$

$$C \text{ \& } A = 4\%$$



Ques.: Genetic Map on basis of foll. R.F.

- QR = 10%
- PQ = 2%
- RS = 8%
- PR = 12%
- QS = 18%



(E) LINKAGE GROUPS

* A linkage group represent a group of linked Genes i.e a chromosome

$$\text{No. of linkage Grp} = \text{no. of chromosome (n)}$$

- eg:-
- Virus + pheno type = 1
 - Neurospora (2n=14) = 7
 - Garden pea (2n=14) = 7
 - wheat (6N=42) = 7
 - Maize (2n=20) = 10
 - Drosophila → 4 (2n=8)
 - ♂ Drosophila = 5 ($\frac{3A+1}{2} \times 1$)

(F) Rate of c.o

\propto dist.

\propto length of chromosome

\propto X-rays (some mutagens)

$\propto \frac{1}{\text{Age}}$

$\propto \frac{1}{\text{Interference}}$

* Interference :- c.o at 1 locus inhibits the c.o of nearby loci.

* Recon = 1 cistron \rightarrow structural gene

Q. c.o in Neurospora

* Neurospora is imp. for study of c.o because it produce ordered tetrad

Q. In a population of Neurospora

100 Ascii are Parental

100 " " non-parental

200 " " Tetra types.

what is F.R.

Sol.

$$R.F = \frac{\text{No. of recombinant} \times 100}{\text{Total Ascii}}$$

$$= \frac{NPD + \frac{1}{2}(TT)}{\text{Total Ascii}}$$

$$= \frac{100 + 100 \times 100}{400}$$

$$= 50\%$$

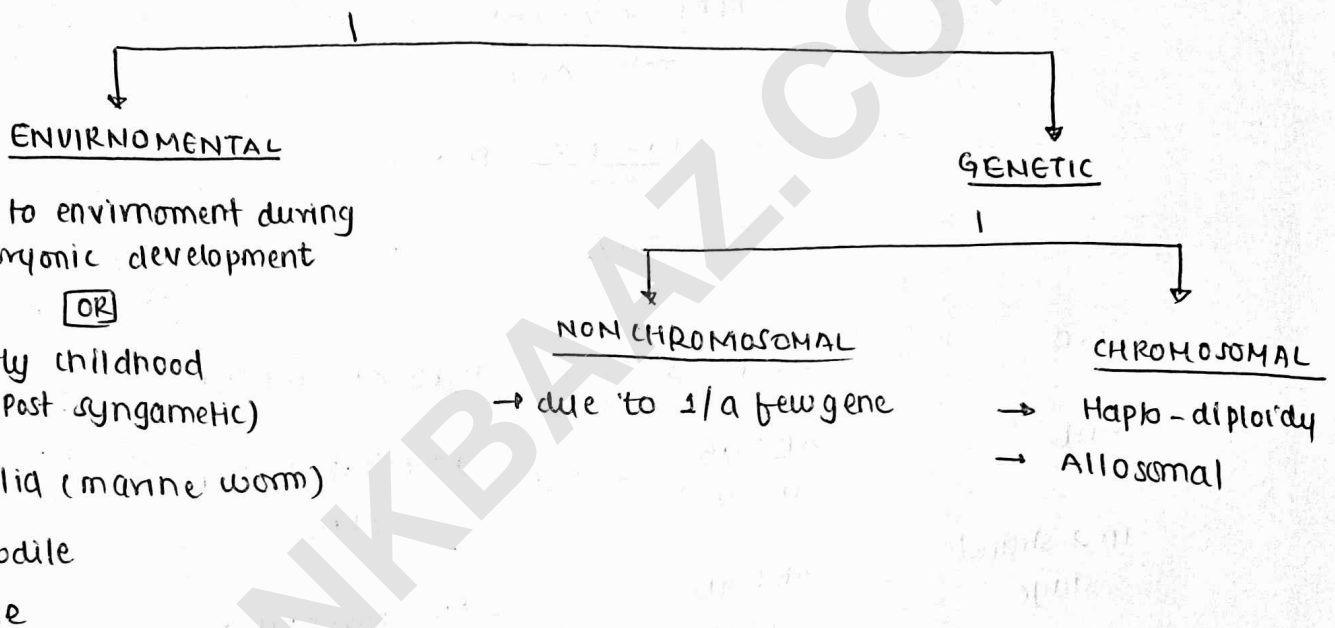
c.o	Ascospores	Type of tetrad
-nt	AB: ab 4 4	Parental Di type (PD)
in 2 strand stage	Ab: aB 4: 4 100% recomb.	Non parental Di type (NPB)
in 4 strand stage	AB: Ab : aB: ab 2: 2 : 2 : 2 50%	Tetra types.

85/ TL-85
 (H) data of linkage in Drosophila

character	Body color	color of eye	shape/size of wings
x-chromosome			
Dominant Trait (wild)	Brown (y^+)	Red (w^+)	Normal (pn^+)
Recessive trait (mutant)	Yellow (y)	white (w)	Miniature (m)

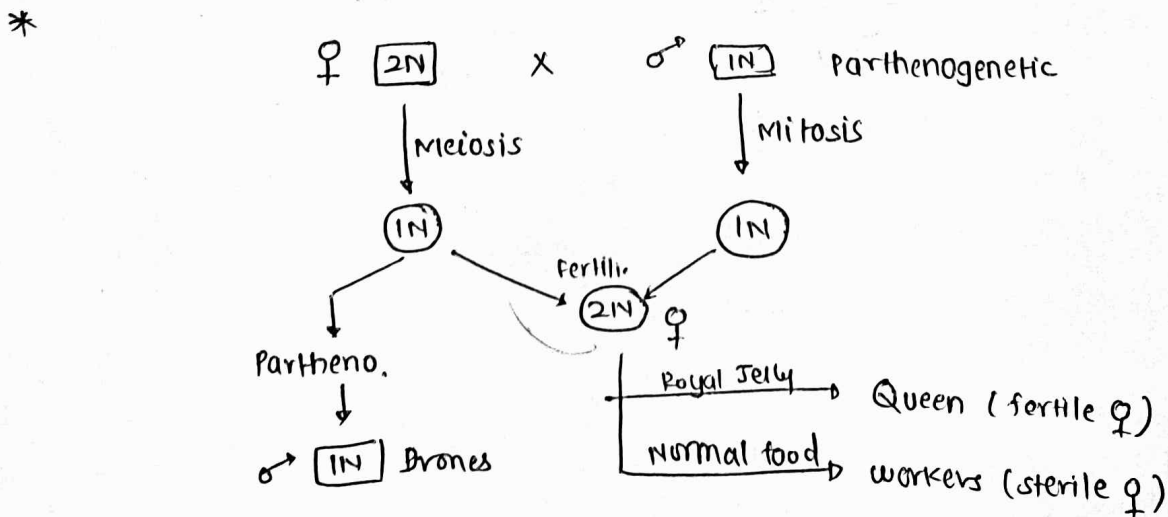
* Male Drosophila always show complete linkage (also for autosomal linkage)

(9) SEX DETERMINATION



(I) HAPLO-DIPLOIDY

eg:- Honey bee ($2N=32$)



* in animals gametes are produced by meiosis & haploid cells do not divide except- parthenogenetic animals.

* Drones do not have father as well as sons however they have grand maternal father & grandson.

II ALLOSOMAL SEX DETERMINATION

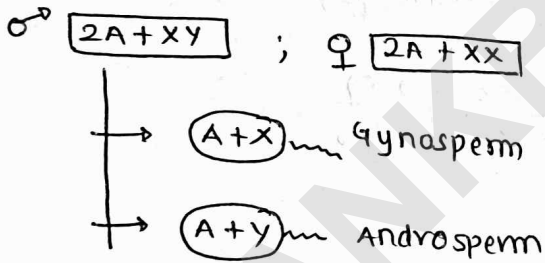
- * Henking :- discovered X-Body
- * McClung :- X-body is X-chromosome + Imp. in sex determination
- * Stevens :- discovery of "Y" chromosome. + its role in S.D.
- * Wilson + Stevens :- "chromosomal theory of sex determination" due to Allosomes.
- * Comens :- "Theory of Heterogametes"

↳ Heterogametic ($2A + XY$) individual is responsible for S.D.
 * Allosomal is syngametic i.e. depends on the syngamy / Type of zygote

♂ Heterogametic
 Allosomes = X & Y

1) XX - XY

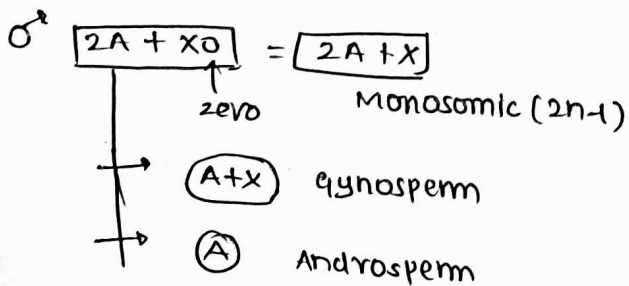
eg:- Humans, *Drosophila**



A = one set of Autosomes

2) XX - XO

eg:- Grasshopper, *Ascaris*



♀ Heterogametic
 Allosomes = Z & W

3) ZZ - ZW

♀ $2A + ZW$

♂ $2A + ZZ$

eg:- Birds, some Reptiles
 some fishes.

4) ZZ - ZO

♀ $2A + ZO$

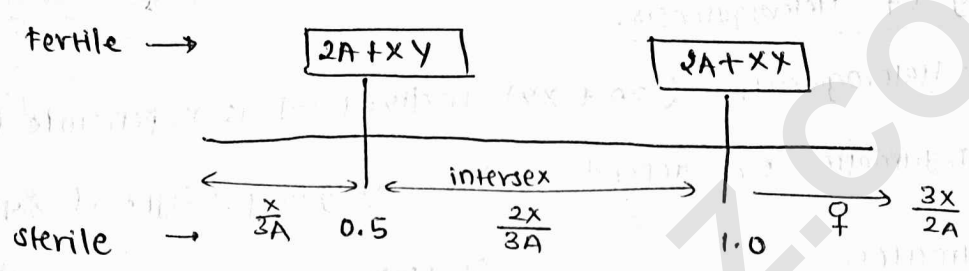
♂ $2A + ZZ$

Rare eg:- some Moth

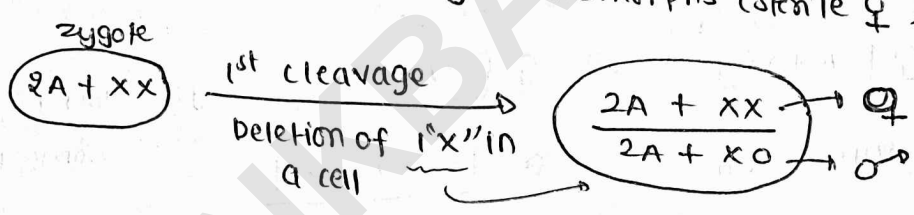
* sex determination in Drosophila

- * Genic - balance Theory by C-bridges (1927)
- * In Drosophila Y-chromosome has no role in sex determination However Y is essential for fertility of male.
- * In Drosophila sex determination depend on sex ratio

$$\text{Genic /sex ratio} = \frac{\text{No. of X}}{\text{No. of 'A'}}$$



* Best evidence - by ~~Eupand~~ Eynandromorphs (sterile ♀)



(10) HUMAN GENETICS

(A) PEDIGREE ANALYSIS

* It is a record of family history + it is important to study the pattern of inheritance of a particular character.

* Mostly it is based on monohybrid cross.

* foll. preference order should be used to solve Q.

(I) Y-linked → All sons are affected if father is affected

(II) Recessive ^{OR} Dominant → Healthy parents but progeny is affected

(III) x-linked ^{OR} Autosomal

6/2
(1) GENETIC DISORDERS

This disorder may ~~auto~~^{chromo}somal or Mendelian.

(A) MENDALIAN DISORDERS

This disorder are due to mutation in 1 or more gene.

(a) SICKLE CELL ANAEMIA

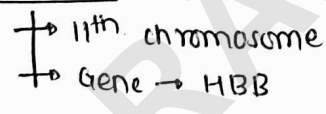
- Gene of β -chain at 11th chromosome
- Autosomal recessive disease However it show codominance at molecular level.
- It is due to substitution of 6th A.A of β -chain of Hb. (Glutamic acid is substituted by valine)
- $Hb^S Hb^S$ in early childhood (sub lethality)
 $Hb^A Hb^S$ R.B.c are normal or biconcave until oxygen conc. is reduced in low O_2 conc. valine forms interactⁿ with other Amino acid which cause reduced permeability of memb., reduced O_2 carrying capacity but more stickyness (Pleiotropic effect)

this individual show more resistance against Malaria.

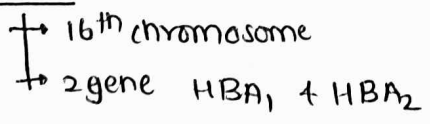
(b) THALLASAEMIA

- it is quantitative or polygenic inheritance which occur due to absence of reduced no. of chain.

for β chain ($\bar{\alpha}$)



For α -chain



- due to frame shift mutation.

HARDY-WEINBERG'S LAW

Q. In a population of Mirabilis red flowered plant are 100, pink flowered 100 & white flowered 200 what is % of R

Sol.

RR Red + Rr Pink + rr white.
 ↓ ↓ ↓
 100 100 200

RR → 100
 Rr → 100
 rr → 200
 R = $\frac{100 + 100}{400} \times 100$

$$\% \text{ of R} = \frac{\text{No. of R}}{\text{Total Allele}} \times 100$$

$$\rightarrow \frac{300}{400} \times 100$$

Q. In a population of Mirabilis white flowered are 64%. What is % of pink & red flowered.

Sol.

$$R+r = 100\%$$

$$\boxed{R+r = 1}$$

$$(R+r)^2 = 1^2$$

⇒

$$\boxed{RR + 2Rr + rr = 1}$$

in ques.

$$rr = 0.64$$

$$r = 0.8$$

$$R = 0.2$$

$$\text{Red} \rightarrow 0.04 \rightarrow 4\%$$

~~White~~ →

$$\text{Pink} \rightarrow 2Rr \rightarrow 2 \times 0.2 \times 0.8 = 32\%$$

Q. In a pop. of garden pea pure tall plants are 300, Hybrid tall 100 & dwarf 100 what is % of hybrid tall plants in 3rd generation

*

$$\begin{array}{ccc} TT + Tt + tt \\ 300 \quad 100 \quad 100 \end{array}$$

$$T = \frac{100}{100} = 0.7$$

$$t = 0.3$$

$$2Tr = 2(0.7)(0.3) = 42\%$$

$$\begin{array}{l} TT = T^2 = 60\% \\ Tt = 2Tt = 42\% \\ tt = t^2 = 20\% \end{array}$$

* Acc. to it Allelic freq. remain const. for generation to generation

- Population size is large.
- there is random mating.
- No evolution. (Natural selectⁿ, mutation should be -nt).
- any process which change allelic freq. should be -nt (Genetic drift, Migration, sampling error should be -nt.)

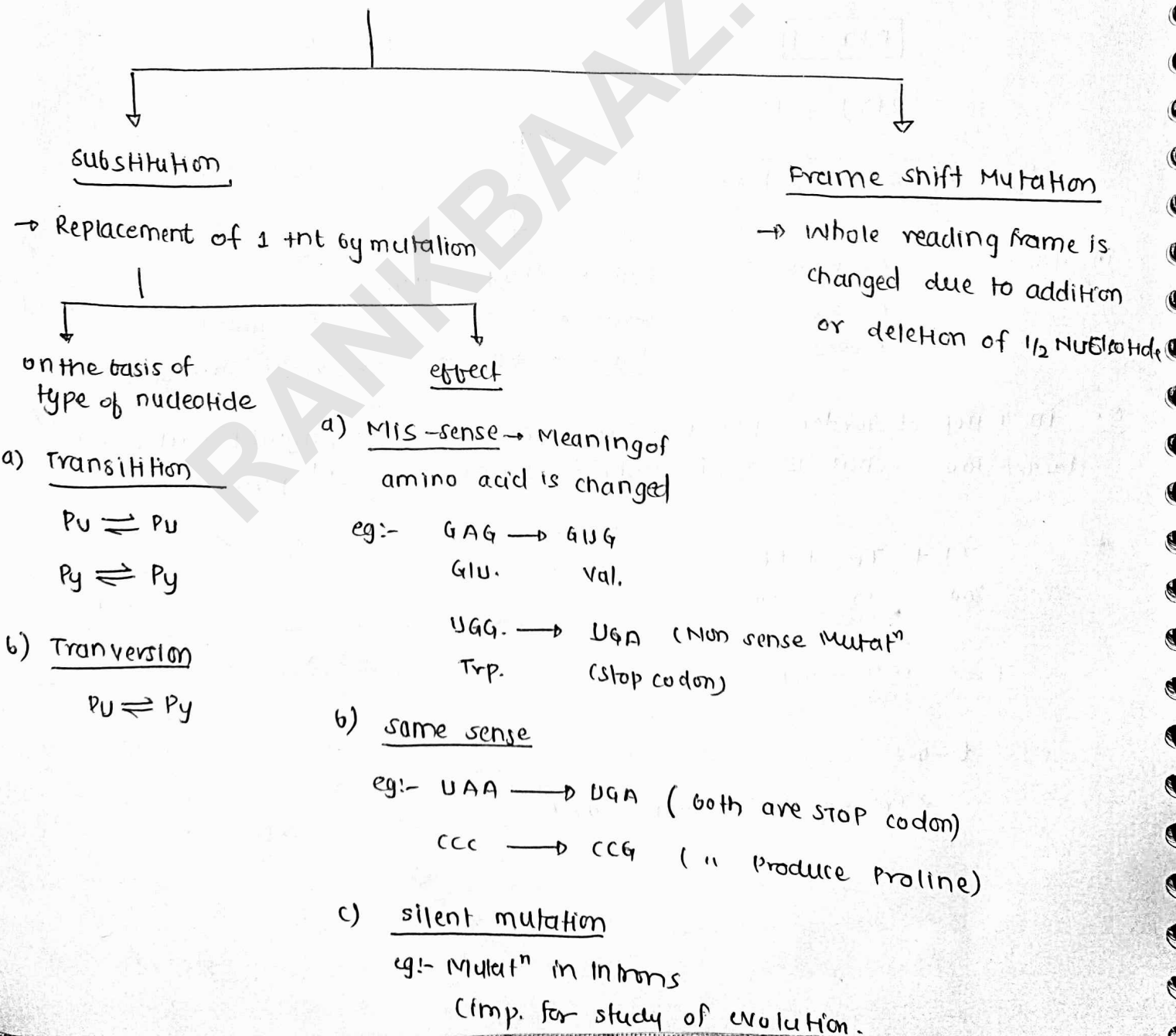
7/02

② MUTATION

* There are sudden changes in genetic material which are inheritable, uncertain non predictable + usually recessive.

* Mutation may occur at gene level or chromosome level.

A) Gene level / DNA level = Point mutation



Substitution

→ Replacement of 1 nt by mutation

on the basis of type of nucleotide

effect

Frame shift Mutation

→ whole reading frame is changed due to addition or deletion of 1/2 nucleotide

a) Missense → Meaning of amino acid is changed

eg:- GAG → GUG
Glu. Val.

UGG → UGA (Non sense Mutatⁿ)
Trp. (Stop codon)

b) same sense

eg:- UAA → UGA (both are STOP codon)
CCC → CCG (" produce Proline)

c) silent mutation

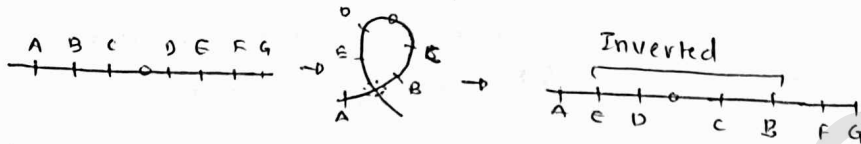
eg:- Mutatⁿ in Introns
(Imp. for study of evolution.)

(B) Mutations at chromosome level

(I) CHANGES IN STRUCTURE OF CHROMOSOMES

- * Para centric \rightarrow without centromere
- * Peri centric \rightarrow involving centromere

- Deletion
- Addition/Insertion
- Inversion



IV) Translocatⁿ (btw Non Homologous chromosome)

- unidirectional
- Reciprocal / bidirectional

\hookrightarrow exchange of gene / chromosome fragments btw non homologous chromosomes

(II) CHANGES IN CHROMOSOME NO.

Aneuploidy

$(2x \pm 1 \text{ or more chromosome})$

Euploidy

* Due to non-disjunctⁿ during meiosis (usually from mother)

Hypoauploidy

(A) Monosomy

$[2x-1]$ ABC
AB-

(B) Nullisomy AB-

$(2x-2)$ AB-
 \hookrightarrow lethal

(C) Double monosomy

$(2x-1-1)$ AB-
A-C

Hyperaneuploidy

(A) Trisomy $(2x+1)$

ABC
ABCe

(B) Tetrasomy $(2x+2)$

(C) Double Trisomy
AABB₂CC₂

Mixed

$2x-1+1$

eg: ABC
A-CC

Q. pollen has 10 chromosome what is no. of monosomic + Trisomic plants.

Ans. 19 + 21 resp.

Q. what is no. of chromosomes in foll. cond.

	Garden Peg ($2N=14$)	wheat ($6N=42$)
Monoploidy ($1N$)	7	7
Haploid ($1x$)	7	21
Diploid Endosperm ($3x$)	21	63
Monosomic ($2x-1$)	13	41
Trisomic ($2x+1$)	15	43

(B) EUPLOIDY

$2N \pm 1N$ or "More N"

* changes in no. of set chromosomes.

i) Autoploidy:- in same species

It may be due to

(a) Parthenogenesis:- $2N [AA] \rightarrow 1N [A]$

'A' = $1N$ of 'A' species

'B' = $1N$ of 'B' species

(b) endomitosis / colchicine treatment

$2N [BB] \rightarrow 4N [BBBB]$

(c) Fertilizatⁿ b/w Apomitic gametes

eg: $2N (\text{egg}) \times 1N (\text{O}^{\text{p}}) \rightarrow 3N (\text{ccc})$ eg. Banana embryo

• Allo euploidy :- due to interspecific hybridizatⁿ

eg:- Raphanobrassica $4N [AABB]$

* **Triticale** $6N$ [AA BB DD]
 $8N$ [AA BB CCDD]

(ii) Autoalloploidy

$6N = AAAABB$

15/2

EXAMPLE OF INTERSPECIFIC HYBRIDS

① Raphanobrassica [$4N = AABB$]

↳ 1st man made / Artificial interspecific hybrid

Raphanus sativus (radish) \times Brassica oleracea (cabbage)
 $2N = 16$ [AA] $2N = 16$ [BB]

"A" = 1N of A species

"B" = 1N of B species

↓
1N

↓
1N

Allopolyploid

(2N)

"AB"

Amphimonodiploid

↓ (b) lchaine

Allotetraploid

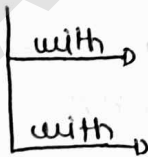
(4N "AABB")

Amphidiploid

② **Triticale** ($6N/8N$) → Man made

secale cereale (Rye)

$2N = 14$ (DD)



Triticum aestivum

($6N = AABBCC$) ⇒ $8N = AABBCCDD = 56$

Triticale

T. durum

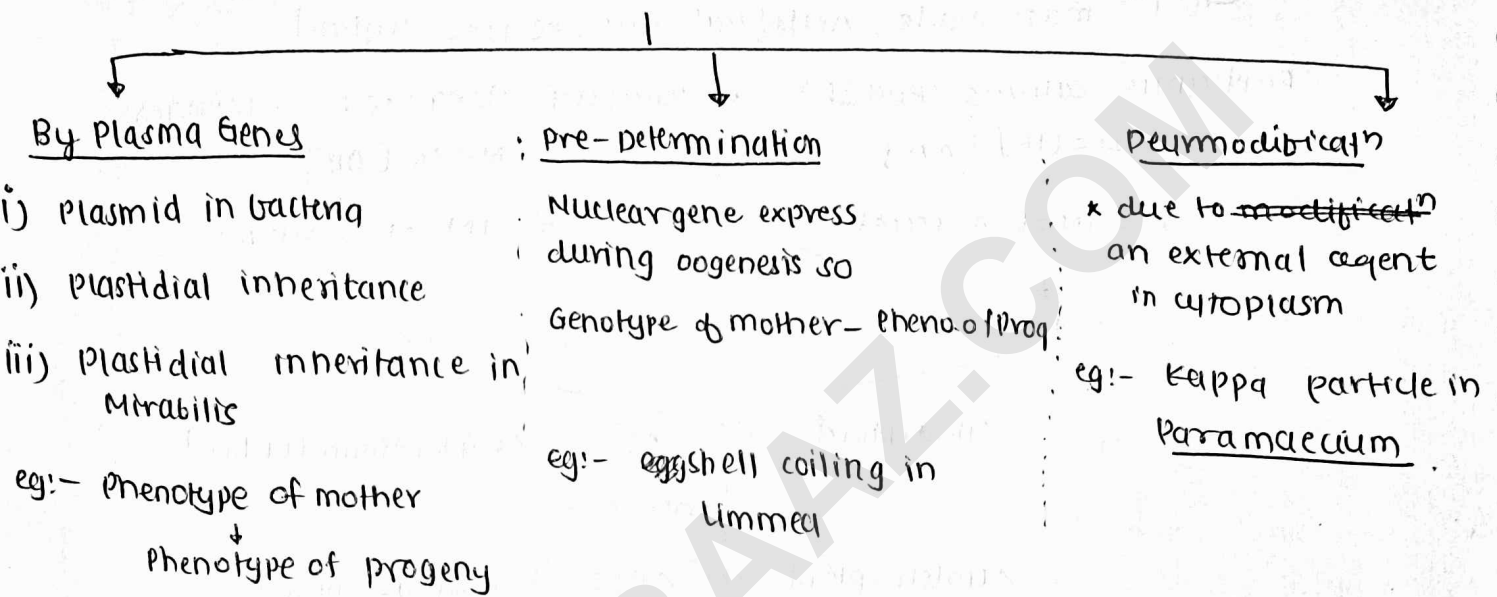
($4N = AABB$) ⇒ $6N = AABBCC = 42$

Triticale

Triticum aestivum } are Natural interspecific hybrid
 durum }

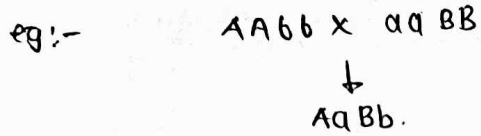
⑬ CYTOPLASMIC INHERITANCE

- * inheritance which is controlled by cytoplasmic genes (Plasma genes) or influenced by cytoplasm
- * it is also called maternal inheritance bcoz zygote ^{receives} cytoplasm from mother parent, it was discovered by cornelis in mirabilis.
- * it is of 3 types.



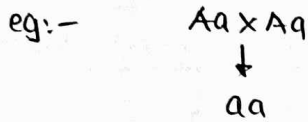
* some other points

- ① sex linked characters → characters controlled by x or y-linked genes.
characters may be somatic or sexual.
- ② sex limited characters → Genes are int on autosomes and are related with 2° sexual characters
(Autosomal)
 - expressⁿ of gene depend on sex hormones.
- ③ sex influenced characters → Expressⁿ of Hetero zygous individual depends on sex hormones
(Autosomal)
 - eg:- Baldness
 - BB → baldness in both ♀ & ♂
 - Bb → No baldness
 - Bb → baldness in male only.
- ④ Hybrid vigour / Heterosis → Hybrid of dissimilar parents show superior characters due to Heterozygous conditⁿ or suppression of recessive alleles
eg:-



⑤ Hybrid breakdown / inbreeding depressⁿ →

⇒ Hybrid of very similar parents or offspring of self pollination may show inferior characters. due to Homozygous condition of recessive alleles.



⑥ Eugenics

* Birth of superior individual is promoted (+ve approach) + birth of inferior individual is inhibited (-ve approach) to improve quality of human population.