

Independent assortment

none of the genes should be linked. They should move freely.

But later understood, the genes in the same chromosome is linked.

Frequency of Recombination measures the intensity of linkage.

At any 1 pt, the

Process of exchange involves only 2 of the 4 chromatids in the meiotic tetrad.

late in prophase I, crossovers become visible as chiasmata.

Non-disjunction → leads to chromosomal irregularities.

Diploid & Tetraploid - stable

Triploid - Not stable.

linked genes can be mapped on a chromosome by studying how often their alleles recombine.

Formation of Chiasmata in late prophase.

Recombination between genes on opposite sides of the crossover pt.

Triple crossover - brings back the original one.

$$\frac{\text{Average no. of crossover}}{0 \times \left(\frac{70}{100}\right) + 1 \times \left(\frac{20}{100}\right) + 2 \times \left(\frac{5}{100}\right) + 3 \times \left(\frac{2}{100}\right)} = 0.42$$

$$\text{Recombination frequency} = 0.3.$$

Recombination

—————
Total # of offspring.

Recombination mapping

with a pt Test cross

Vg Vg BB

R.F btw Vg & b is 18%

This is equal to (18) map

units / 100 CM on the

genetic map.

No recombinants - Recombinants.

$$0 \times 0.82 + (1) \times 0.18 = 0.18$$

Vg⁺ b⁺

Vg⁺ b⁺

F₁ = Vg

Vg b

Vg b

415

405

92

88

Frequency of recombination =

Problem

Maize

leaves = 2 col. alleles

G → purple leaves (D)

g → green " (R)

S → short stalk (R)

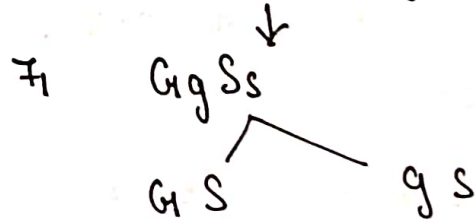
s → Tall " (D)

F₁ → purple & Tall.

When they were back crossed to plants with

g & s, produced an F₂ in which.

G G S S × g g s s



F₁ ⇒ G g S s × g g s s

Total 200 plants.

- (1) green short (2) purple tall
- (3) greentall (4) purple short

(1) ⇒ what is the evidence that

New combination is possible only if u have recombination

Parental no (high)

R. Com (No less)

1 cross over (NO ↓)
2 cross over (NO ↓)

Total 200

Parental

75, 79

Recombinants =

24, 22

BT

we calculate R.F

Total no. of Recombinants

offspring

0.3 is less than 0.5

RF

Cross overing happens

Repulsion heterozygote

Crossing over does not

coupling heterozygote.

0.23 is the RF

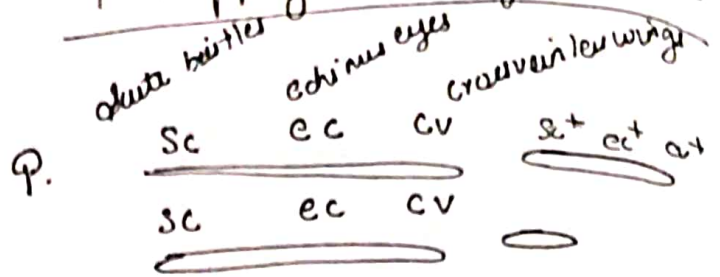
BT

Genetic distance is directly

proportional to R.F.

0.23 Morgan

R. Mapping with 3pt Testcross



light side by side align. just opposite.

Pseudoautosomal region.

for segregate properly.

Pair with one region.



Beginning - 2 types of chromosomes

↓
8 diff chromosomes.

| | | | |
|----------------------------|-----------------|-----------------|-----------------|
| dute, echinus, nonvirious. | Sc | ec | cv |
| wild type | sc ⁺ | ec ⁺ | cv ⁺ |
| dute | sc | ec ⁺ | cv ⁺ |
| E, cross | sc ⁺ | ec | cv |
| Scv, ech | sc | ec | cv ⁺ |
| cross | sc ⁺ | ec ⁺ | cv |
| Scv, cross | sc | ec ⁺ | cv ⁺ |
| Echinus | sc ⁺ | ec | cv ⁺ |

Extensions of Mendelism

Incomplete dominance
 Co
 Multiple alleles
 Epistasis
 pleiotropy
 In breeding

(Find out cog of incomplete & co dominance)

Mendel's works

diff alleles may affect phenotype in diff ways
 # one gene - 2 alleles more than
 # Genes \rightarrow 2 alleles

IGT \rightarrow produce epitope.

Incomplete Dominance 1:2:1

WW = Red 1 (more protein)
 Ww = Pink 2 (less pro)
 ww = white 1 (loss / change of function)

In heterozygous condition dominant allele can't fully mask the recessive allele
 Intermediate phenotype.
 one allele is partially / incompletely dominant over the other

Yellow + Red \rightarrow orange
 ↓
 Intermediate phenotype

change of function
 Code for one protein
 Code for another protein

Co-dominance

No intermediate phenotype.
 Combination of both
 No merging
 Have both dominant
 1 : 2 : 1 Ratio.

Multiple Alleles

- ① $c^+c^+ \rightarrow$ WT
- ② $c^h c^h$ - white hair with black tips on the body
- ③ $ch ch \rightarrow$ black hair
- ④ $cc \rightarrow$ white hair.

Hierarchical phenotype.
 suppressed / dominant
 The most common \rightarrow WT
 All others are mutant.

For 1 loci \rightarrow diff phenotype
 When multiple alleles \rightarrow dominant allele will be in hierarchical

one gene controlling many

P phenotype.

For a given loci you

can have 2 alleles

more 2 alleles for

F phenotype.

Blood group also

Single gene is controlling

but have diff phenotype.

F multiple alleles → also an example of Codominance.

Allelic series.

hypomorphic alleles →

partial function

A null allele → non

functional

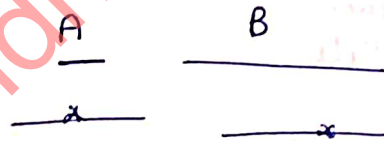
An allelic series describes dominance hierarchy of multiple alleles

F

When 2 or more characters are inherited, individual hereditary factors assort independently ^{during} gamete production, giving diff traits → Equal opportunity of occurring together.

During gamete formation, the alleles for each gene segregate from each other so that gamete carry only one allele for each gene.

mutations
Testing for allelism.



Same phenotype.

only in homozygous condi

(Recessive)

Understand

white gene.

white locus + white eye phenoty

Caused by several genes.

How eye colour of Dros is controlled by several genes?

Heterosis

F

2 diff in bred line
crossed, the hybrids are
heterogous.

Genetic analysis of
inbreeding

TtRr x TtRr

↓

Tall Round
Short wrinkled

new parent
Tall wrinkled
Short Round

3:1

↑

new combination is produced

Genetics of eye colour in Drosophila - Assignment

linkage / crossing over /
Recombination.

When two are
present in chromosome
?

They will segregate
together.

Independent assortment
will not happen.

All the genes present in one
particular chromosome in.

You will see some sort of
variation

When two genes present in non-
homologous chromosome, they segregate
independently with 100% efficiency.

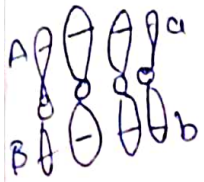
They show any reduction in the
chromosome.

linkage \Rightarrow 2 genes are linked
together.

When 2 genes are linked,
they segregate together, Ab.
independent assortment.

Beyond meiosis & mitosis -
S phase - doubling happens

- Tetrad - 4 -



homologous pair.



crossing over



chromatids

sister chromatids.

Recombination - second step in evolution.

More variation.

Recombination can happen through independent assortment / segregation.

Genes that are on the same chromosome travel through meiosis together

Thomas Hunt Morgan +

Wild

Mutant

$b^+ b$

bb

$vg^+ vg$

$vg vg$

Test cross.

Ratio = 1:1

$b^+ vg^+$ Homozygous

$b vg$ homozygous recessive

TTSS $vt+ss = 1:1:1:1$

$b^+ b$

$b b$

$vg^+ vg$

$vg vg$

Both are in the same chromosome.

that combination were absent in the chromosome.

Does not only.

Just pair and into family potential.

Can not hold be placed on the chromosome even if placed on the chromosome → Recombination.

Wants = 100%

Frequency = 0%

Frequency - 50% or 0.5