

An Institute for IAS Exam
(By a Group of JNUites)
Web: www.l2a.in Email: l2adelhi@gmail.com

Anthropology (Test code: AN02TS09-18)

4-19

Mobile no: 9899776680

Email: ani.ruddh981@gmail.com

Name of the Candidate ANI RUDDH YADAV

CSE Admit card no

Place ORN

Time 11:15

Test No.

4

Class room

Distance Learning

Date

06/07/2019

INDEX TABLE

Q. No.	Maximum Marks	Marks Obtained
1	50	25
2	50	
3	50	29.5
4	50	30
5	50	29
6	50	29
7	50	
8	50	
9		
10		
11		
12		
13		
14		
15		
16		
17		
Total Marks Obtained		142.5

INSTRUCTIONS

1. Do furnish the appropriate details in the answer sheet (viz., Name, Admit card no and Test Code).
2. There are **EIGHT** questions printed both in English and Hindi.
3. Candidates has to attempt **FIVE** questions in all.
4. Question No. 1 and 5 are compulsory and out of the remaining, **THREE** are to be attempted choosing at least ONE from each section.
5. The number of marks carried by a question/part is indicated against it.
6. Answers must be written in the medium authorized in the Admission Certificate, which must be stated clearly on the cover of this Question-cum-Answer Booklet in the space provided. No marks will be given for answers written in medium other than the authorized one.
7. Word limit in questions, wherever specified, should be adhered to.
8. Answers of questions shall be counted in chronological order.
9. Any page or portion left blank in the Question-cum-Answer Booklet must be clearly struck off.

Signature of Examiner

L2A™

Feedback/Comments

Do not
write
anything in
this margin

91-4-1

Dear Aniruddh.

attempted well.

L2A™

Do not
write
anything in
this margin

→ 40 points related to evolving
significant points

Anthropology Test Series -2018

AN02TS09- 18

Time allowed: **Three Hours**

Maximum Marks: 250

Candidates has to attempt **FIVE** questions in all. Question No. **1** and **5** are compulsory and out of the remaining, **THREE** are to be attempted choosing at least **ONE** from each section. The number of marks carried by a question/part is indicated against it.

Word limit in questions, wherever specified, should be adhered to.

Attempts of questions shall be counted in chronological order. Unless struck off, attempt of a question shall be counted even if attempted partly. Any page or portion left blank in the answer book must be clearly stuck off.

This is Question-cum-Answer booklet, questions to be attempted in the provided space.

SECTION - A

Q1. Write notes on the following in about 150 words each

10x5=50

10

(a) Problems and prospects of Tribal development in North-East India

Adaptive Radiation

Adaptive radiation is the process of descent from a common ancestor where evolutionary process gives multiple descents with varying phenotypes with respect to one or many traits.

Features

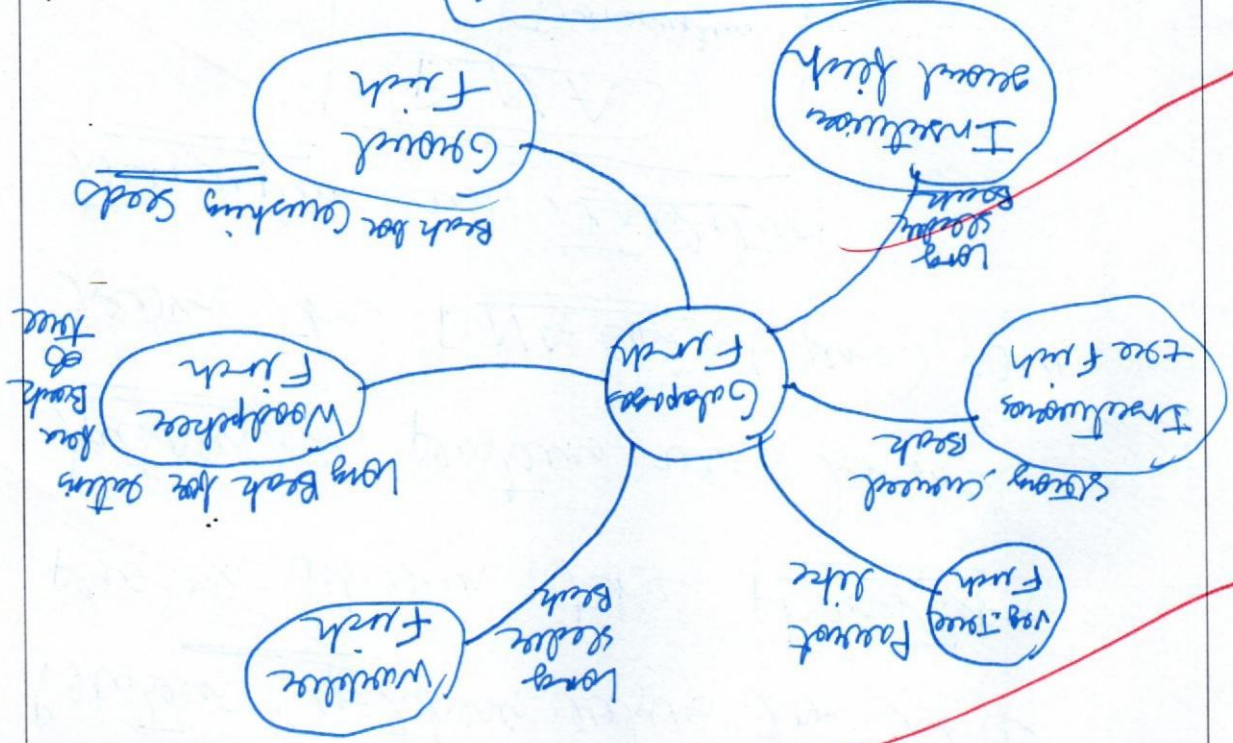
It occurs due to various reasons

- ① Increased competition in an area due to similar niche
- ② Isolation and Speciation
- ③ Development of fresh environmental pressures

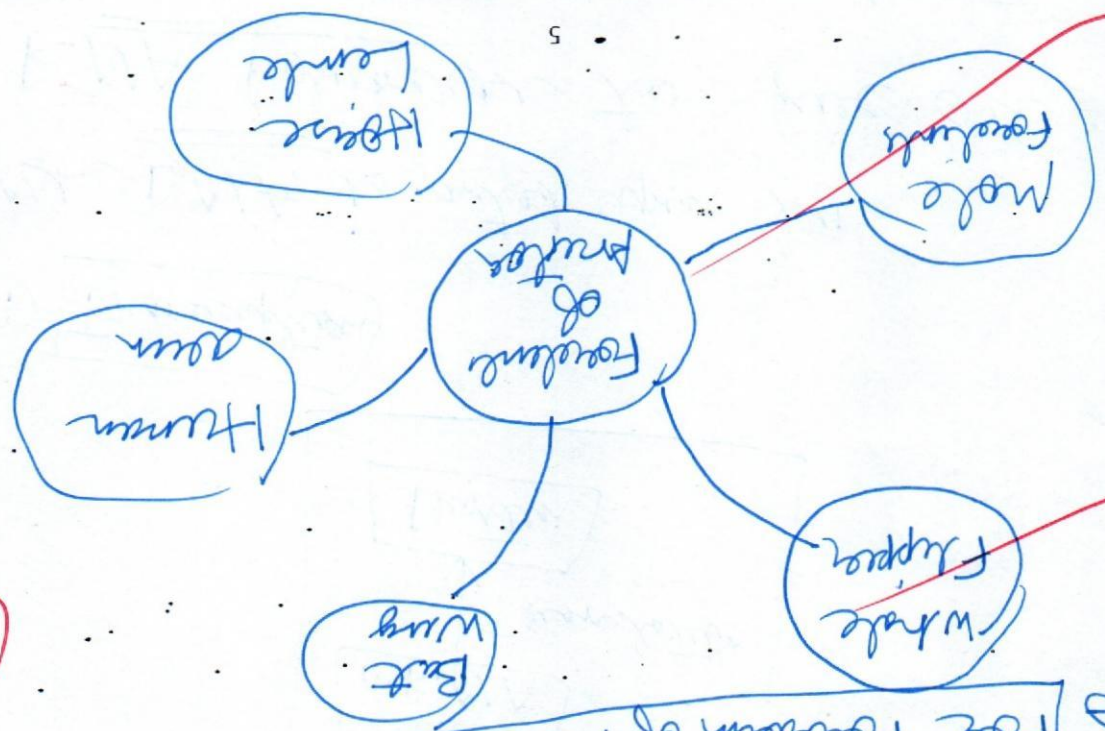
L2A

Eg = C. Darwin studied the evolution of finches of Galapagos Islands and gave the following for each structure

Do not write anything in this margin



→ For Evolution of Mammals

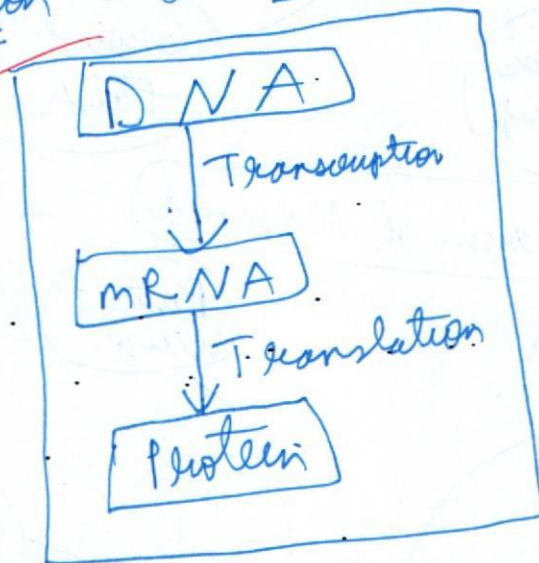


6

(b) Positive discrimination

Protein Synthesis

Protein Synthesis refers to the process wherein the phenotypic factors or proteins are formed from the DNA code (gene) via transcription and translation.

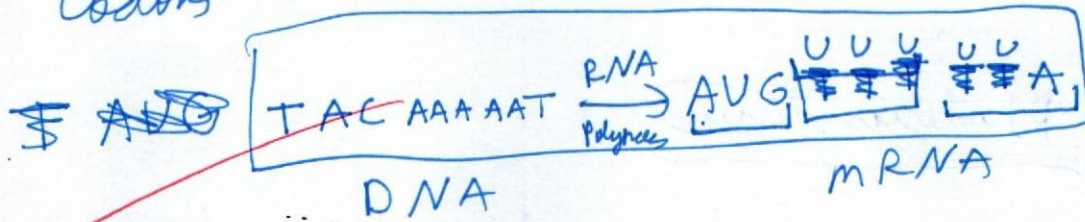
① Transcription

The DNA is acted upon by RNA Polymerase to give an mRNA Transcript containing information.

of Amino Acids in the form of
Codons

L2A™

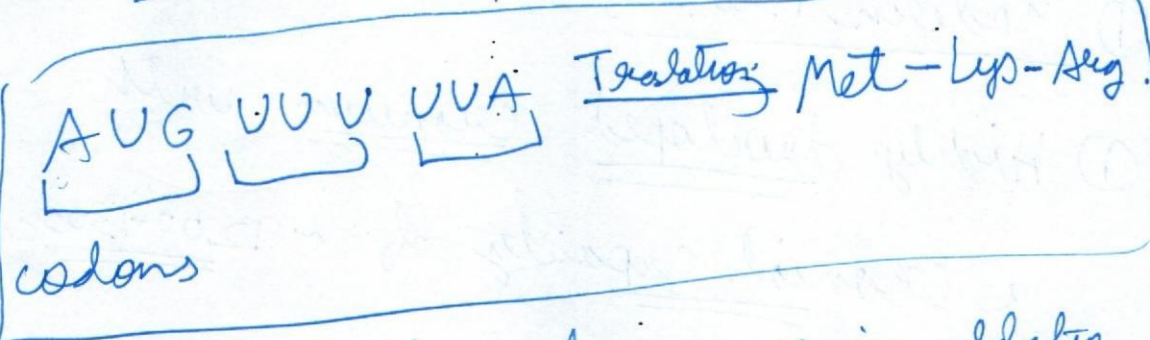
Do not
write
anything in
this margin



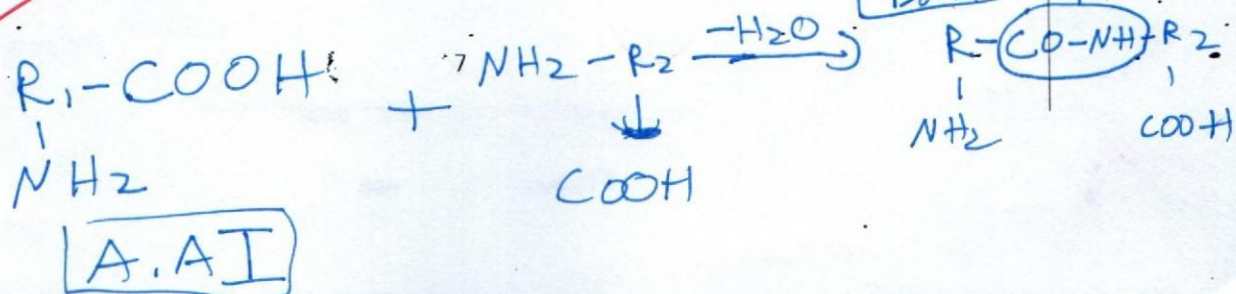
Translation

The mRNA transcript is then transported
to the cytoplasm via the
nucleus for translation using the
ribosomes on tRNA adaptor

Complex



This happens due to successive addition
of Amino Acids via peptide Bond



(c) Linguistic Anthropol Bio-genetic variability of Indian tribes

Rhodesian Man

Rhodesian Man refers to a fossil of a skull found in Zambia representing a ~1.5 - 1 million year old fossil that is anachronistic with its appearance.

Features① Modern Features

① Highly developed cranium with a cranial capacity of ~1200-1300cc

②



Revise

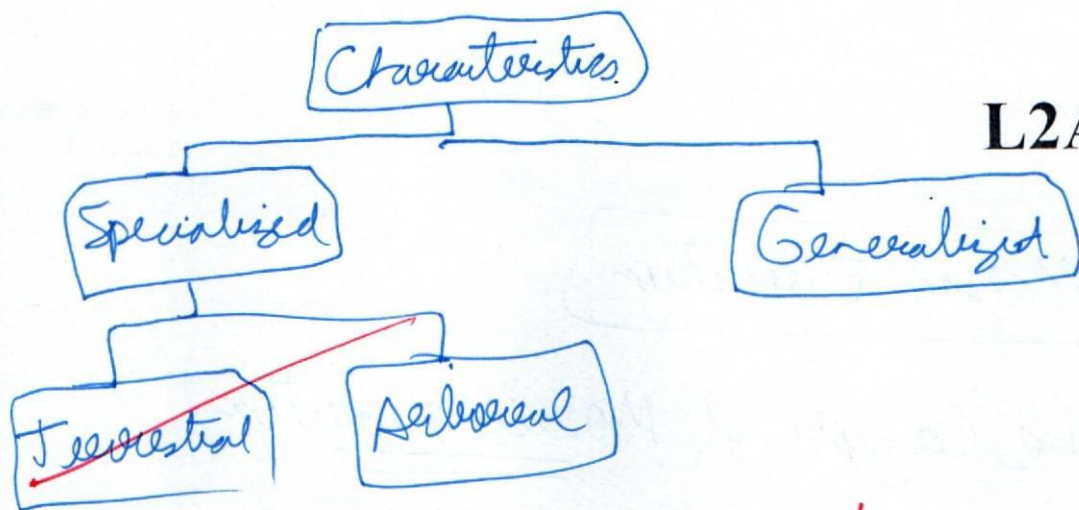
(d) Pseudo-tribalism

PrimatesDefinition

Mivart (1873) has defined primates
as ungulate, clavicate, placental
mammals with at least 3 types
of teeth at one point of life;
well developed brain and forebrain;
developed extremities with at least
one pair of opposable digits;
Hallux with developed nail or
none; scrotum testes, penis
pendulous, pectoral mammellae.

Characteristics

Characteristics of ¹⁰ primates can be of
two types



L2A™

Do not
write
anything in
this margin

General Characters

- ① Developed forelimbs of rachis,
- ~~② Ultra etc~~
- ② Development of opposable digit
for grasping
- ~~③ Brain Capacity developed due to
larger Cerebrum~~
- ④ Penis Pendulous
- ~~⑤ Pectoral Mammal~~
- ~~⑥ Sacroter Sac-like~~

6
54.
7 1/2

(e) Starvation Deaths and PVTG's

Mosaic Evolution

The Concept of Mosaic evolution was proposed by Rutherford who said that evolution occurs in a gradual manner in an unsymmetrical and non-equitable manner.

eg.

Evolution Form	Time (million Yrs Ago)	Encephalization	Bipedalization
Homo Erectus	2 MYA	800-900cc	Biped
Homo Neanderthalensis	0.3	~1500cc	Biped
Homo Sapiens	.04	~1400cc	Biped, Fully
Australopithecine	3-2 MYA	~600cc	Partially Biped
Homo Habilis	2-1.5 MYA	~700cc	Partially Biped

Thus from the temporal perspective
the encephalization and bipedalization
of man has evolved in an
independent and seemingly unrelated
manner however both have had
a significant impact on the general
and cultural evolution of man.

eg. High degree of encephalization
made head bigger thus higher
time of infant dependency, thus
led to male-female exclusive bond

Thus different traits show a mosaic
pattern in evolution however all
work in a manner to ensure
overall evolution

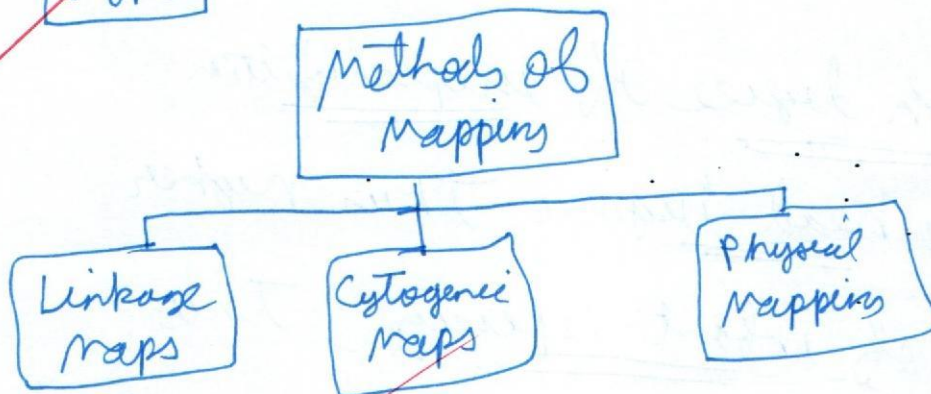
Q2 (a) What is the concept of ethnicity? Provide explanations of Ethnic Violence between the Upper Caste Hindus and Scheduled Castes

20

Gene Mapping

Gene Mapping refers to the process of linking genes to specific regions or loci on the genome as that provides an overall contextual map of gene placement on chromosomes.

Types

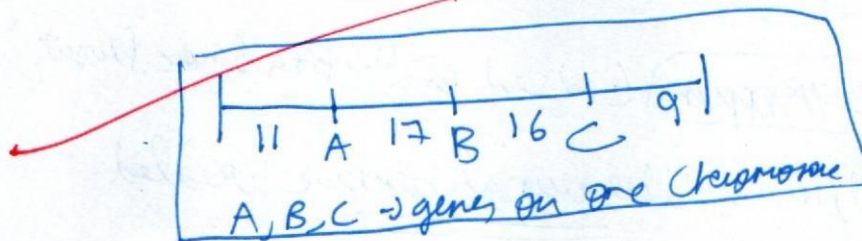


① Linkage Mapping

It occurs when multiple genes on one chromosome are linked through hybrid experiments. Eg. A dihybrid

product is used for a double cross
experiment or trihybrid for ~~triple~~ **L2ATM**
~~cross~~ quadruple cross experiment.

Do not
write
anything in
this margin



② Cytogenetic Maps

It is prepared by tracing location
of specific genes on specific chromosomes

① Hybridisation

Gene present on specific chromosome is
amplified using hybridisation primers
in a PCR to confirm presence

① In Situ Hybridisation

Using a labelled probe to match
on a particular gene to detect
location on chromosome by FISH

③ Physical Maps

15

① Restriction Mapping

Restriction Mapping involves using restriction enzymes to fragment DNA and consequently map specific genes to different loci

L2A™

Do not
write
anything in
this margin

(b) Contiguous Mapping (Used in Human Genome Project)

→ It involves a library of genome created using restriction enzymes, consequent sequencing using shotgun sequencing, identifying and mapping genes based on contiguous fragments.

Human Genome Study

The Human genome was first fully sequenced by the Human Genome Project under Dr. Craig Venter and the results were:-

- ① 3 Billion Base pairs of DNA
- ② 99.99% DNA is common to all
- ③ Africa contributed as the cradle of civilization

④ 27000 - 32000 genes in human

⑤ 95% 'Junk DNA'

Thus the Human genome project

L2A™

15

(b) Write a note on the linguistic distribution of Indian Tribal population.

Do not
write
anything in
this margin

provides us with a lot of information
for our genetic basis of life
and allowed us to study genes,
sequences etc.

Applications

① Disease prediction through genome
analysis or sequencing

② Directed gene therapy and customised
medicine

③ Genome study has caused loss of
global identity of all races
despite differences

④ Use in Environment and Agriculture which
uses gene information for better environment
and better crops

⑤ Forensic Anthropology ⑥ Paternity Analysis.

12/2

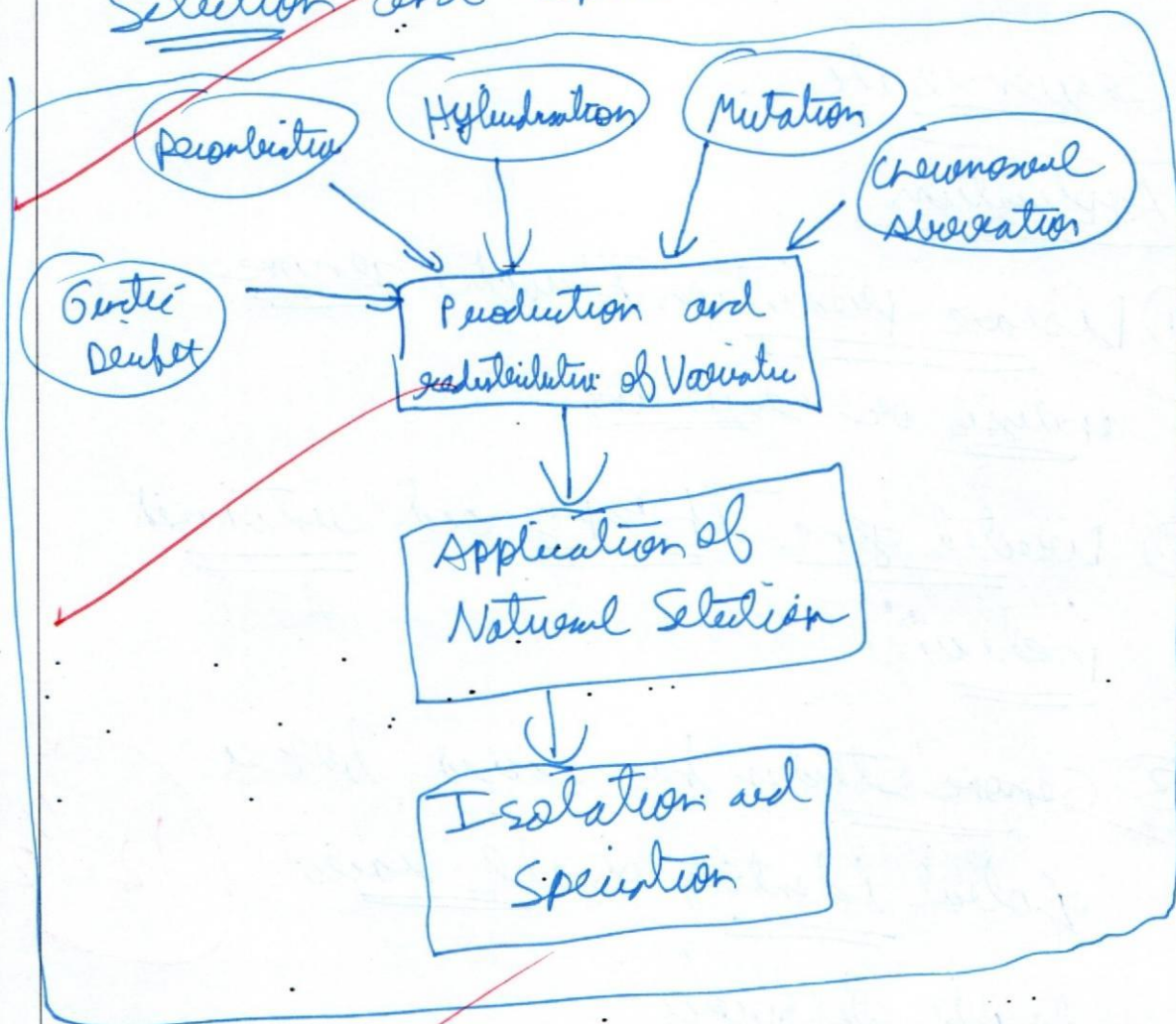
Q2 (b)

Evolution as described by the Synthetic Theory of evolution

L2A™

Do not
write
anything in
this margin

is a combination of multiple
processes of mutation, natural
selection and speciation



Thus variation / mutation and natural
selection can be understood as the
two drivers of evolution.

Effect of Mutation

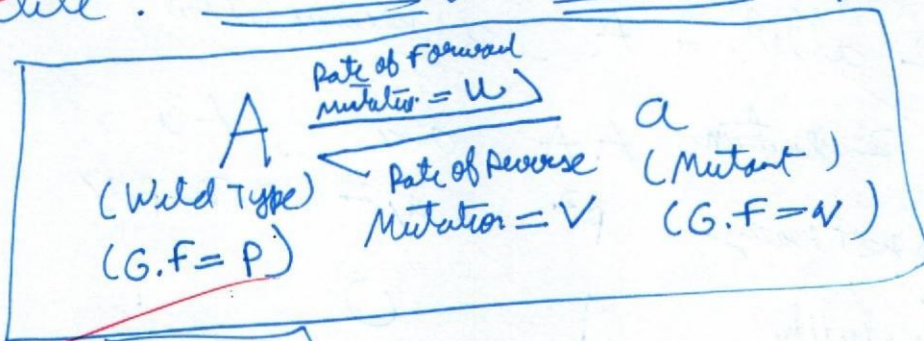
Mutation refers to any spontaneous change in DNA and if germinal then heritable

L2A™

Do not
write
anything in
this margin

Mathematical proof of mutation effect

Let 'A' and 'a' be 2 alleles in a population with 'a' as the mutant allele. Gene frequency is p and q respectively



$$p + q = 1$$

Change of Gene frequency of population $\Rightarrow p u - v q = \Delta q$

In Hardy Weinberg Equilibrium; $\Delta q = 0$

$$p u = v q ; p + q = 1$$

$$p = \frac{v}{u+v}$$

$$q = \frac{u}{u+v}$$

Thus mutation is affected by rate of mutation in a population only in process of evolution

Natural Selection

Refers to the process through which

L2A™

(c) Critically evaluate reservation policies for SCs and STs and to what extent they have led to social mobility among them.

15

Do not write anything in this margin

a selection mechanism ensures viability of reproductive success in a given environment.

Ex. Natural Selection is a recessive lethal allele

Initial population \Rightarrow ~~AA~~ \rightarrow P

\Rightarrow 2 alleles \Rightarrow A, a (recessive lethal)

\therefore 2 genotypes	AA	aa	Aa
Gen frequency	p^2	q^2	$2pq$
Viability	1	0	1

Thus \Rightarrow the total population $= (p^2 \times 1) + (2pq \times 1) + (q^2 \times 0)$

$$= p^2 + 2pq$$

$$= p(p + 2q)$$

$$= p(1 + q)$$

also add
dynamic

equilibrium between
M & N.S.

(8/12) Δ in gene frequency of 'a' $\Rightarrow \Delta q = q - \frac{1}{2} \times \frac{2pq}{p(1+q)}$

$$\Delta q = \frac{q^2}{1+q}$$

\therefore Natural Selection is inefficient in weeding out recessive lethal allele which depends on initial gene frequency only

Genetic load

Genetic load refers to the relative
decrease in fitness of a population
due to deviation of the genetic
bases from the ideal optimum
genetic situation.

$L \rightarrow$ Genetic Load

$I \rightarrow$ Ideal / optimum

$w \rightarrow$ Fitness of population

$$L = I - w$$

Significance of Genetic load

- ① It signifies a deviation from
the ideal / optimum genetic fitness
of the population which represents
a 'load'

Depends
by any
factor -
muller etc.

① It signifies a sort of 'genetic death' of individuals due to reduced fitness of individuals which consequently causes a reduced reproductive success rate

L2A™

Do not
write
anything in
this margin

Types of Load

① Mutational Load

It refers to the decrease in fitness due to mutations in population

↳ (a) Silent Mutation → No effect on load

(b) Missense Mutation → Slight effect on load

(c) Nonsense Mutation → Significant effect on load
frame shift

② Indicative Load or Genetic Load

Some situations do not allow

the gamete to be viable

or survival proceeding this leads to
death eg Tetrisomy of chromosomes!

L2A™

Q3. (a) Discuss the religious conversions in Tribal India with suitable examples.

15

Do not
write
anything in
this margin

Factors that abet load

- ① Mutation: Rate of Mutation will abet genetic load
- ② Migration: Inbreed of population with different genotypes will abet the load
- ③ Selection: Varying amounts of selection due to changed environment

Limitation

84%

- ① Excess 'genetic load' causes extinction
- ② It is not necessarily a negative phenomenon as it only reduces fitness with reference to a particular environment

849

Definition

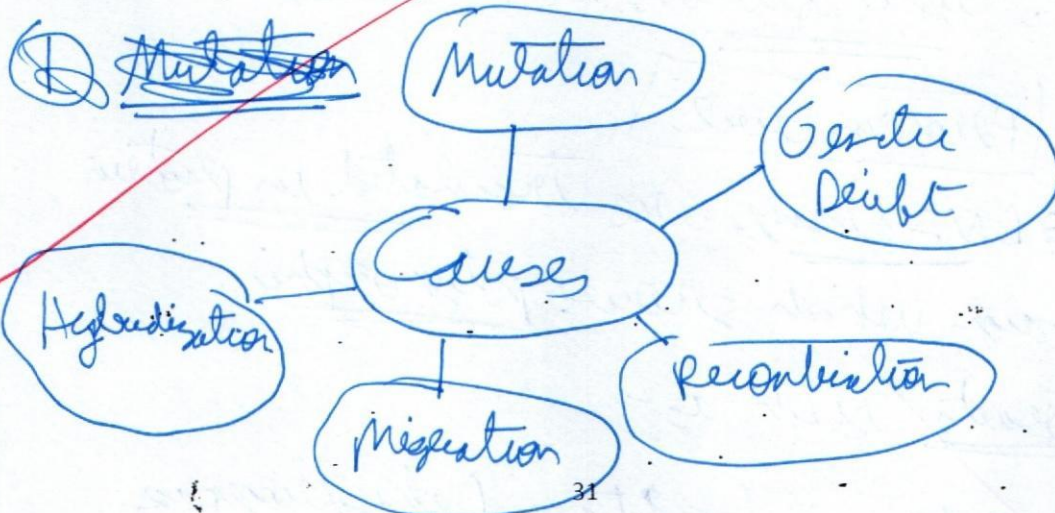
L2A™

Do not
write
anything in
this margin

Genetic Polymorphism refers to ~~the~~ the occurrence of two or more distinct forms of alleles such that the rarest of them cannot be maintained by mutation alone.

It is only considered a polymorphism if it occurs in more than 1% of the population.

Cause of Genetic Polymorphism



Types of Genetic Polymorphism Seen

① Cell Surface level

L2A™

Q4 (a) Given historical analysis of forestry in British and Post-British India.

20

Do not
write
anything in
this margin

~~eg~~ A, B, O, AB antigens are coded for
by Transferrin on Chromosome 9 (RBC)

① Rh Factor gene → Chromosome 1 (RBC)

② HLA Antigen (WBC)

Thus various polymorphic forms of antigens
present on cell surface

② DNA Level

All changes have underlying effect from
DNA, thus there must be a genetic
change.

eg. Cystic fibrosis, Huntington Disease

③ Protein level

DNA changes are translated in protein
changes which shows polymorphic
traits such as:-

① 5 types of Lactase Dehydrogenase

② Different types of Hemoglobin

such as Chaperone, Hithava etc

L2A™

Do not
write
anything in
this margin

④ Chromosome level

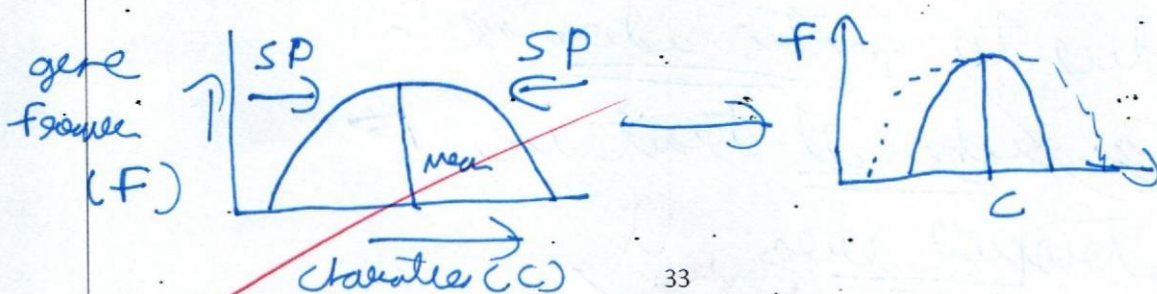
Polymorphic forms of Chromosome represent diseases often eg trisomy of 21 (Down's Syndrome); Structural alteration (Angelman Syndrome) etc.

Evolutionary Forces

① Prevalent Polymorphisms and Stabilizing Selection

Extreme polymorphism are removed by selection which preserves the mean form

eg. Inters cannot reproduce thus
2 species maintain



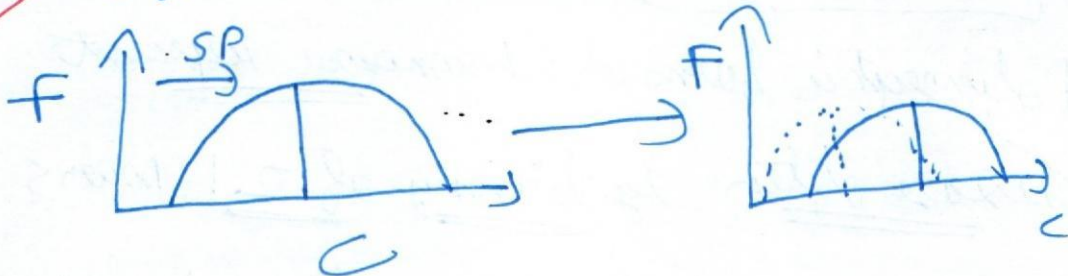
SP → Selection Pressure

② Transient Polymorphism and Directional Flow

One extreme form is favored due to type of environment

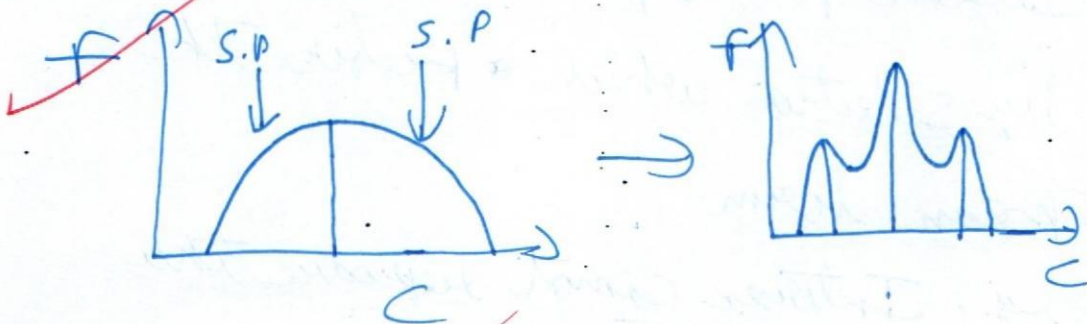
L2A™

Do not write anything in this margin



eg Blood group B in oriental populations

③ Stable polymorphism and Heterozygote advantage



It preserves heterozygous individuals due to genetic advantage,
eg. Sickle cell trait in tropical areas

attempted well

(b) Critically examine the Development Induced to Displacement of Tribals

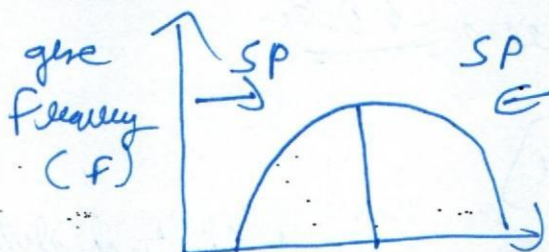
Natural Selection

Natural Selection refers to the set of processes that determine evolutionary change in a population by ensuring maximum genetic viability is chosen and reproductive success is guaranteed in the population.

Types of Selection

① BASED on effort

① Stabilizing Selection



Character

SP → Selection Pressure

35



eg. Intense individuals are selected against to stabilise genes the 2 in humans

say the mechanism to pre-require 7 selection

② Directional Change



L2A™

Do not
write
anything in
this margin

Selection flows to one direction

eg Selection of B ~~for~~ blood group in
particular Oriental population.

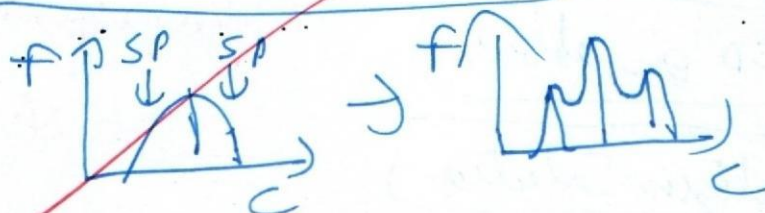
③ Disruptive Selection



Intermediate form is not chosen

eg African Swallow Tail Butterfly can only
survive if it mimics predator

④ Heterozygous Selection



Heterozygous forms are chosen

⑤ Based On Magnitude

Depends on Kin, frequency and density of selective
traits.

⑥ Based on Ecological Traits

① R-Selection → Large population, High resources, favours
low variability and high population

② K-Selection → Small population, Scarce resources, favours
individuals

Hardy Weinberg Law

According to the Hardy Weinberg law "In a Mendelian population the allele frequencies and genotype frequencies remain constant over generations if there is no effect of mutation, migration or selection."

This situation is referred to as Hardy Weinberg Equilibrium wherein allele frequencies and genotype frequencies are constant.

Conditions for Equilibrium

- ① Large population with random Mating
- ② No Mutation is operational
- ③ No Selection is operational
- ④ No Migration or Closed Population

Proof

Assume 2 Genotypes AA , aa

L2A™

with gene frequencies p and q . In the next generation we get Aa ; thus

	$A (p)$	$a (q)$
$A (p)$	$AA (p^2)$	$Aa (pq)$
$a (q)$	$Aa (pq)$	$aa (q^2)$

Genotypes $\rightarrow AA (p^2)$, $Aa (2pq)$, $aa (q^2)$

Thus Hardy Weinberg gave the equation

$$p^2 + 2pq + q^2 = (p+q)^2$$

Gene frequency $\Rightarrow A \rightarrow \frac{p^2 + pq}{p^2 + 2pq + q^2} = \frac{p(p+q)}{(p+q)^2} = \underline{p}$

$$a (p+q=1)$$

$$\Rightarrow a \Rightarrow \frac{q^2 + pq}{p^2 + 2pq + q^2} = \frac{q(q+p)}{(p+q)^2} = q$$

Thus gene frequency and genotype frequency remain conserved over generations.

Limitation of Hardy Weinberg

(1) Purely theoretical concept due to the assumptions of infinitely

Do not write anything in this margin

large population, no Mutation and no Selection

L2A™

Do not
write
anything in
this margin

② ~~There is no way to ~~to~~ combine the~~
~~absence~~ of genotypic mutations unless
phenotypic character is affected

③ Selection is based on the environment
which is dynamic thus cannot be
stopped

④ ~~Only valid for large populations~~
~~in controlled population~~

Significance of Hardy Weinberg

① ~~Reflects a conservative force that~~
~~is to be overcome to ~~to~~ ensure~~
~~evolution.~~

② Reflects the stable nature of
Selection ~~which operates only~~
~~when evolved~~

③ ~~It explains heterozygous advantage~~

8½

SECTION - B

Q5. Write notes on the following in about 150 words each

10x5=50

10

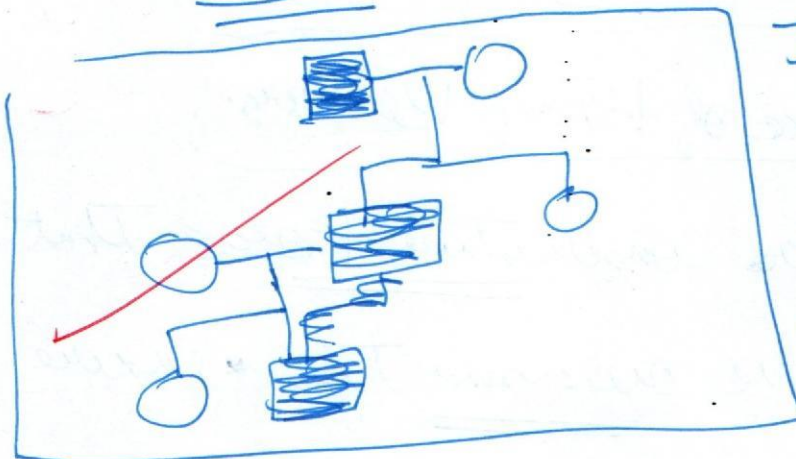
(a) Meaning of Ethnic Identity in the contemporary world

Sex linked Characters

Characters that are linked to the
Sex Chromosome →

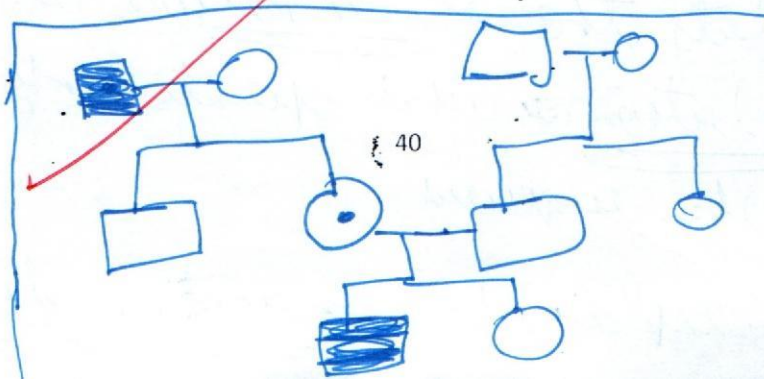
Male → XY
Female → XX

eg. Hairy ears in ~~Men~~ Men is a
Y linked



It only
affects men
of the same
line and all
generations

eg. Hemophilia (X linked Recessive)



Affects
males more
due to
Hemophilia

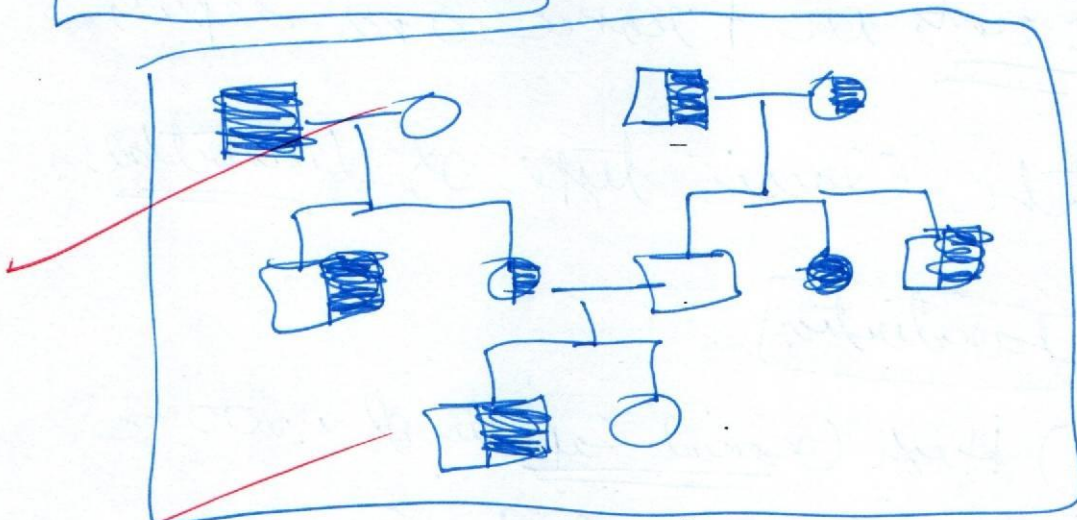
Sex limited Characters

L2A™

Do not
write
anything in
this margin

Characters that are inherited
independent of the sex chromosomes
and are thus autosomal in nature

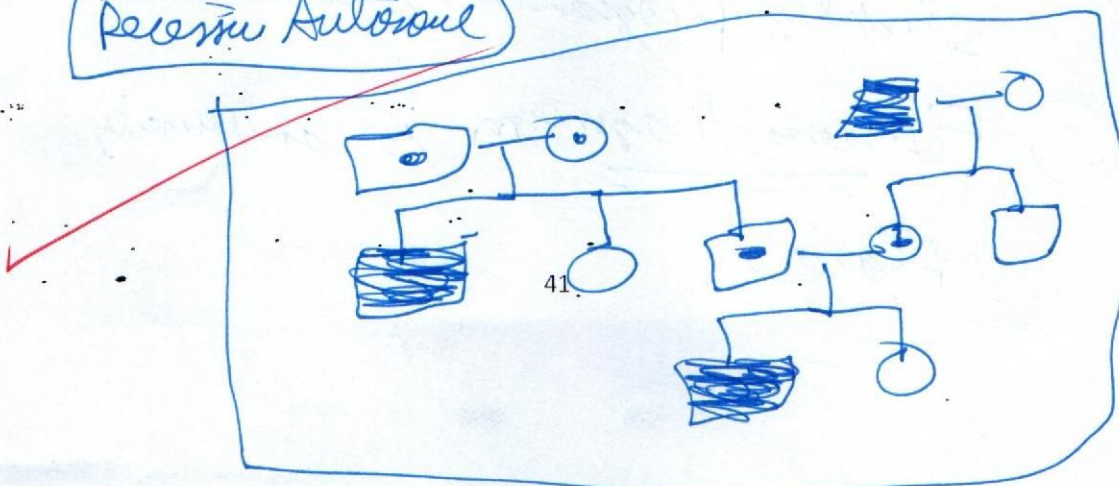
Dominant Autosomal



It affects all genders equally due
to autosomal nature and heterozygosity
is enough for expression

6

Recessive Autosomal



(b) Denotified tribes

Do not
write
anything in
this margin

La Chapelle Aux Saints is a
fossil of Homo Neanderthalensis

discovered in La Chapelle Aux
Saints in France and depicts
the 'Classic' type of Neanderthal.

Characteristics

- ① High Cranial Capacity of ~1600 cc
- ② Rough texture of Skull
- ③ Broad base and wide
- ④ Parabolic dentition with
slightly prognathic features
- ⑤ Foramen Magnum is anteriorly
placed

- ⑥ Developed like Asperu
- ⑦ Very Stout and Broad
- ⑧ The Face structure is not completely human like

The Chapele Ar Saut also had

Technology

Cultural Features

- ① Clothing → Bone Needles Found

- ② Tools → Sophisticated ~~not~~ not lower level tools

- ③ Bone Cult → Advanced Bone Shells found

- ④ Case Histories

6

(c) Special Component Plan & Tribal Sub Plan

Genetic Counselling refers to the process of determining the actual or potential genetic status of a person and informing them of the possible consequences of their genetic predispositions.

Method

Screening



Risk Analysis



Management of
Issue

① Screening → Involves DNA sequencing, RFLP, VNTR, etc. to screen for possible.

disease.

In case of Prenatal mother CVC or
amniocentesis is done

L2A™

Do not
write
anything in
this margin

② Risk Analysis

Polygenic traits are harder to
analyse but monogenic diseases such
as Thalassemia, ~~Sickle~~ Sickle cell
anemia etc. can be detected

③ Management

① Gene Therapy

↳ Embryo Therapy

↳ Gene silencing using siRNA

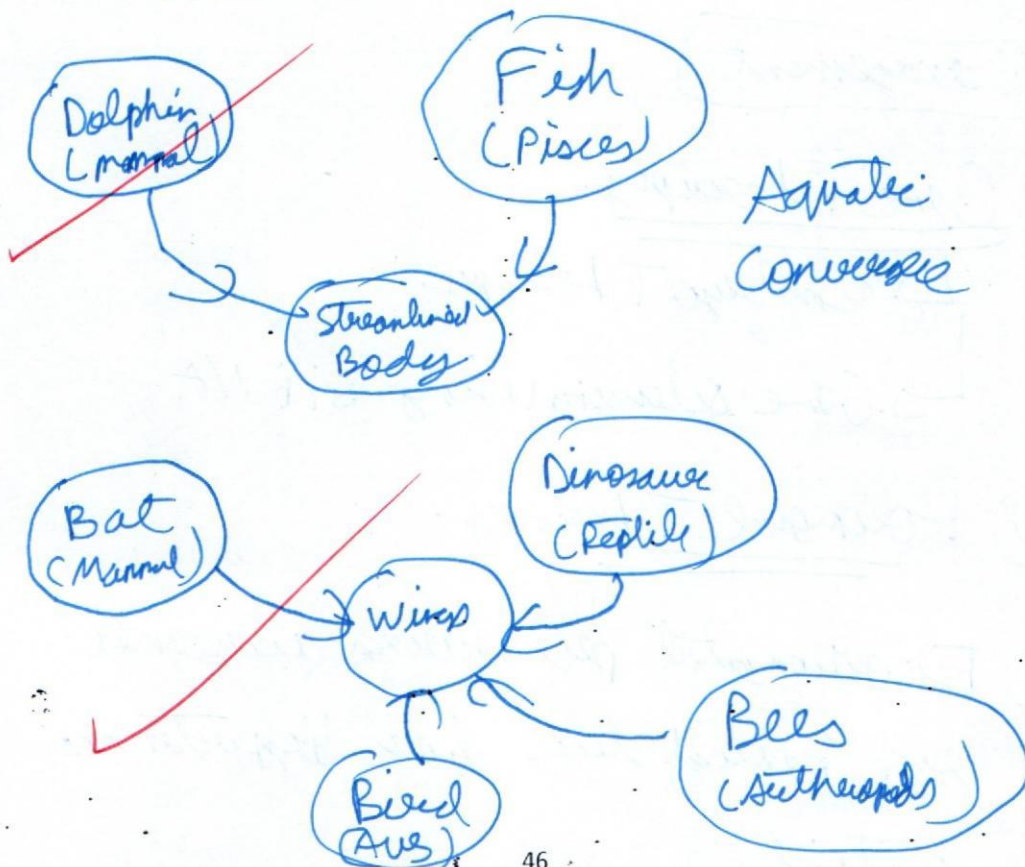
② Hormonal Therapy

③ Environmental pressures awareness
eg. Reduced stress for Hypertension
patient

6

Convergent Evolution

It refers to the convergence of
~~evolutionary~~ features due to
 similar need but does not
show similar structure and only
similar function.



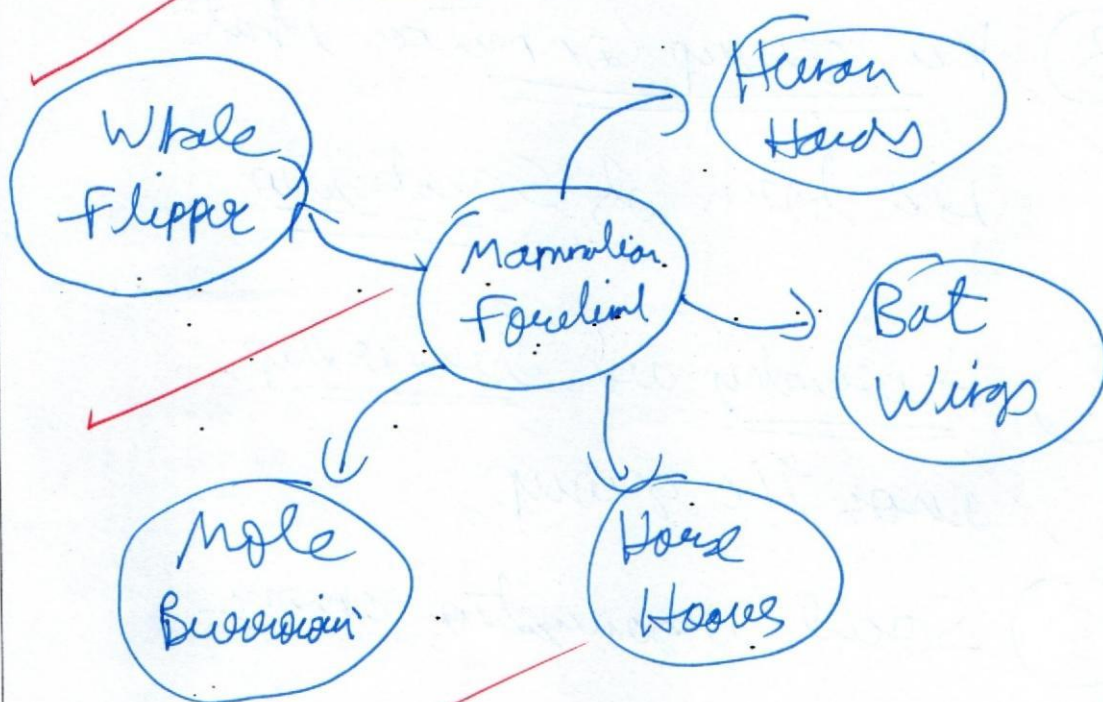
Divergent Evolution

L2ATM

Do not
write
anything in
this margin

It refers to divergence of evolutionary
traits or ~~function~~ that were once
~~shared~~ by the descendants' common
ancestor

They share similar ~~function~~ structure
but ~~not~~ function.



add
retermnant too

B. size
D. size
D. size

L2A™

(c) Relevance of Tribal Panchsheel in the light of emerging development practices

10

Do not
write
anything in
this margin

Primate Social Behaviour

availability
of
resources
predation
etc.

① Formation of Social groups

② Forms of Communication

↳ Olfactory (smell)

↳ Auditory (chest beating, grunting)

③ kin group correlation that
is a form of socialization

④ Hierarchy and leadership
among the group

⑤ Social organization among
groups

⑥ Band like structure among
gorilla.

88a

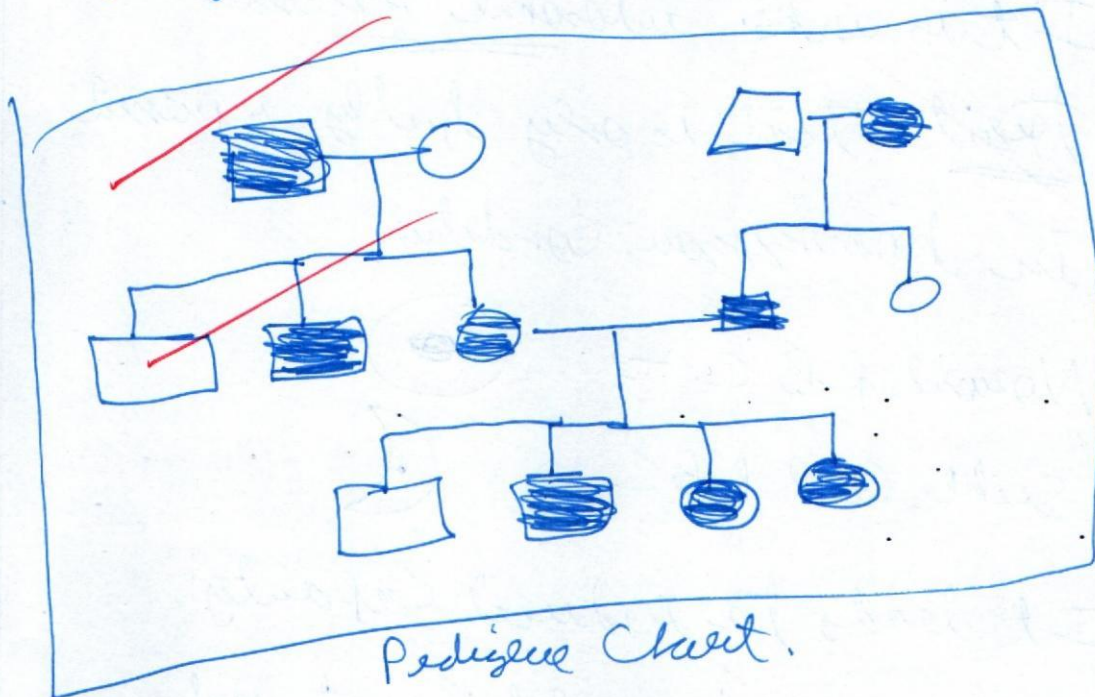
Molecular Basis of 3 inherited disease

L2A™

Do not
write
anything in
this margin

① Huntington's Disease / Disorder

It is an autosomal dominant disease which results in an inherited genetic condition. It can be passed on from both mother or father.



This is caused by one simple genetic polymorphism of DNA and

they can be ⁶⁷ cured by
gene editing / therapy. Early



Q8. (a) Discuss the socio-cultural, economic, and psychological constraints responsible for low literacy in tribal areas. Give your suggestions

15

Diagnosis is possible with
perinatal amniotic sample or
chorionic Villi sampling.

② Sickle Cell Anemia

→ It is an autosomal recessive
trait that is only fully expressed
in homozygous condition

→ Normal RBC → 
Sickle Cell RBC → 

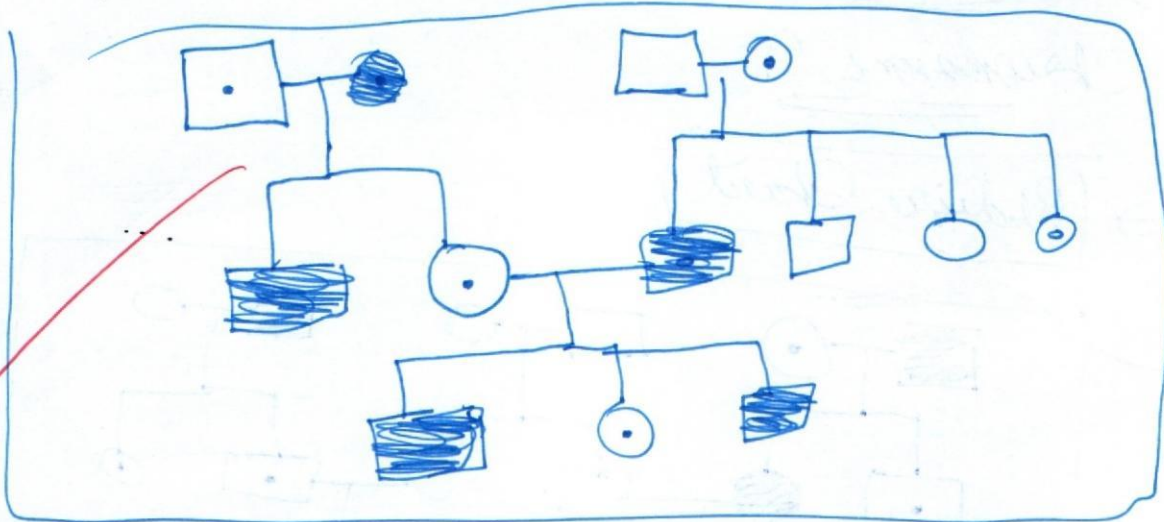
→ It leads to reduced capacity
to carry hemoglobin ~~and~~ and
thus causing tiredness and lack of
health

→ It is caused by ⁶⁸ single polymorphisms
on the gene coding for

Pedigree Chart

L2A™

Do not
write
anything in
this margin



Hemophilia causes a reduced amount
of hemoglobin thus leading
to anemic conditions due to decreased
capacity to carry blood.

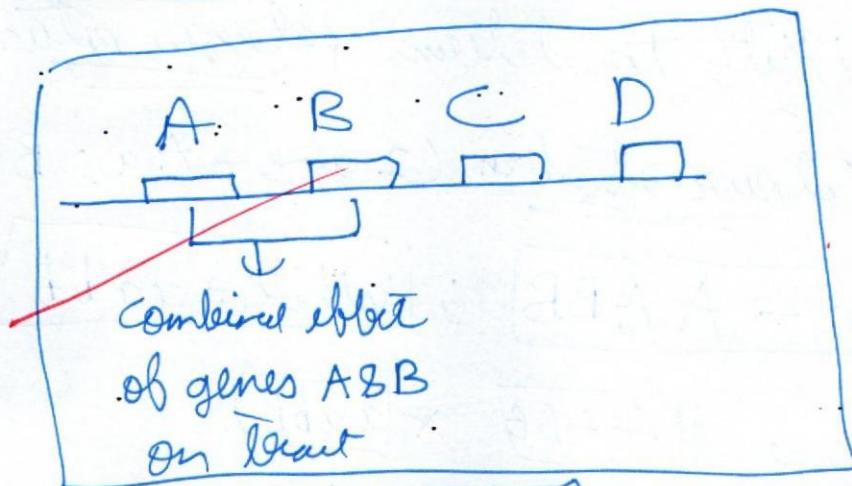
12

(b) Critically analyze the impact of development programmes and welfare measures on tribals and weaker sections.

20

Polygenic Inheritance

Polygenic inheritance refers to the mechanisms of inheritance of certain quantitative traits that are continuous in nature and thus are not affected by one single gene loci but multiple gene loci.



Multifactor Hypothesis

R. H. Fisher gave the hypothesis which states that certain

continuous traits such as height,
intelligence etc are affected by
multiple genetic factors and thus
do not ~~to~~ follow Mendelian pattern
of inheritance with respect of traits
inheritance.

Darwin's Experiment

Darwin used a cross breeding
observation experiment between Niggers
and Whites to determine polygenetic
of colour of skin (2 genes $\rightarrow A, a ; B, b$)

Nigger $\rightarrow A A B B$; White $\rightarrow a a b b$

F_1 cross \rightarrow $A A B B \times a a b b$

F_1 Genotype
 F_1 Phenotype

$A a B b$
72
mulattoes

L2A™

Do not
write
anything in
this margin

F₂ Cross AaBb x AaBb

→ male ↓ Female	AB	Ab	aB	ab
AB	AABB	AABb	AaBB	AaBb
Ab	AABb	AAbb	AaBb	Aabb
aB	AaBB	AaBb	aaBB	aaBb
ab	AaBb	Aabb	aaBb	aabb

L2A™

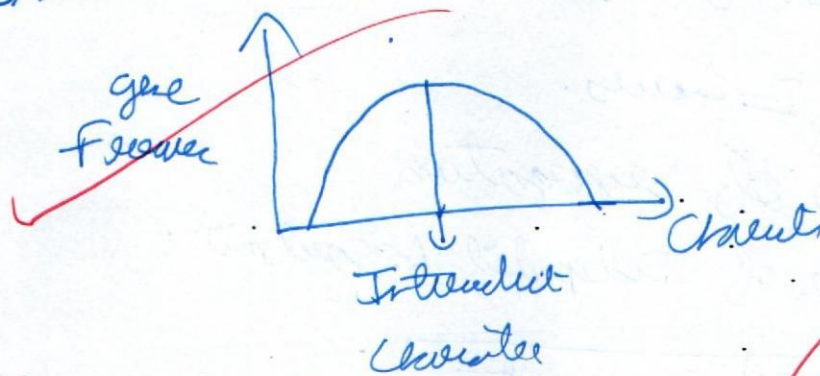
Do not
write
anything in
this margin

∴ Genotypic Ratio → 1 : 4 : 6 : 4 : 1

or Black : Dark : Mottled : Light : White

∴ Quantitative Observation

Distribution of Polygenic traits



(Effects in Man)

- ① Continuous traits → Height, intelligence etc.
- ② Threshold effect → Colour of eyes
- ③ Pleiotropic effect
- ④ ~~Co~~ Dominance
- ⑤ Incomplete Dominance

Mendelian Genetics refers to the branch dealing with inheritance of monogenic traits that follow the Mendelian pattern of inheritance.

Mendelian Pattern of inheritance

Mendel's experiments with pea saturnum were based on 7 characters and thus he gave 2 Laws.

- ① Law of Segregation
- ② Law of Independent Assortment

① Law of Segregation

Each trait or character is determined by a pair of alleles that are inherited independently and during genetic process are segregated separately.

Hand-drawn diagram illustrating a monohybrid cross between a tall (TT) and a dwarf (tt) pea plant. The P generation (TT x tt) produces an F1 generation (Tt x Tt). The F1 generation is then selfed (Tt x Tt) to produce the F2 generation, which consists of four offspring: two tall (Tt) and two dwarf (Tt).

Do not
write
anything in
this margin

~~F_2~~

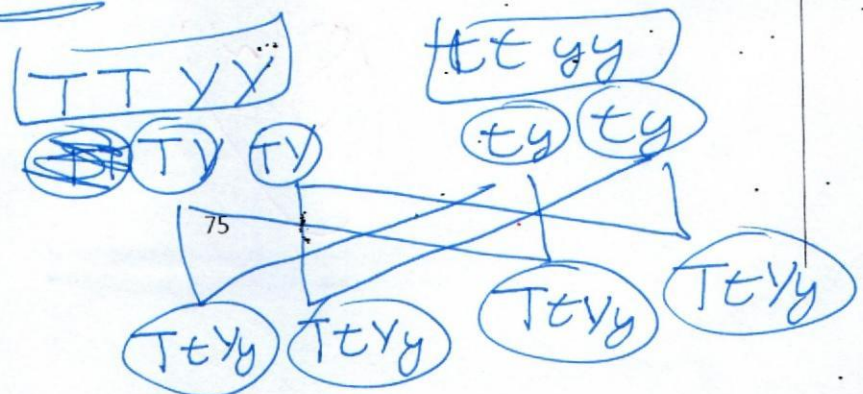
	T	t
T	TT	Tt
t	tT	tt

\therefore There is segregation
of gametes

② Law of Independent Assortment

Multiple factor inheritance occurs in
an ancestral independent manner the
inheritance of a separate factor or
trait.

is, 2 traits are inherited independently of each other.

 ~~$\frac{F_1}{F_2}$~~ 

F ₂	TY	Ty	tY	ty
TY	TTYY	TTYy	TtYY	Ttyy
tY	TtYY	Ttyy	ttYY	ttYy
Ty	TTYy	Ttyy	TtYy	Ttyy
ty	TtYy	Ttyy	ttYy	ttyy

Do not
write
anything in
this margin

~~∴ genotype frequency → 1 : 3 : 6 : 3 : 1~~

∴ Phenotype frequency → $1 : \frac{3}{4} : 9 : \frac{3}{4} : 1$

Mendelian Inheritance in Man

- ① A, B, O Blood group
- ② Genetic diseases such as Huntington's disease
- ③ Sex linked diseases such as Hemophilia etc
- ④ Achondroplasia

gkn