

Anthropology 215

CAUSES OF EVOLUTION



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Summary of Hardy-Weinberg

In the absence of any disrupting factors the allele and genotype frequencies at any given locus in a randomly mating population will be repeated faithfully from one generation to the next; should the frequencies be perturbed for any reason, they will return to the expected equilibrium values after one generation of random mating (Giesel, 1974:14)

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Predictions of H-W

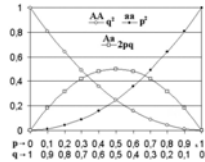
- Predict the offspring's genotypic proportions from the frequency of the alleles present in the parent's generation. For a trait determined by alleles, the proportion of the 3 genotypes is $p^2 : 2pq : q^2$



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Hardy-Weinberg

2. Frequencies of genes (alleles) of a population are inherently stable from one generation to the next. i.e. the proportion p^2 $2pq$ q^2 for a two-allele system will continue generation after generation.
3. In the event that two populations merge, an equilibrium of gene frequencies is reached immediately. The first hybrid population will be in a state of genetic equilibrium.



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The Utility of H-W

- It allows estimates of departures from equilibrium to be measured.
- Can predict genotypic frequencies when gene frequencies are known and random mating is assumed.
- Can calculate the number of heterozygote carriers of autosomal dominant conditions.



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Which population is in Hardy-Weinberg Equilibrium?

	AA	Aa	aa	p	q
I	.20	.80	.00	.6	.4
II	.36	.48	.16	.6	.4
III	.50	.20	.30	.6	.4
IV	.60	.00	.40	.6	.4

Answer

II .36 .48 .16

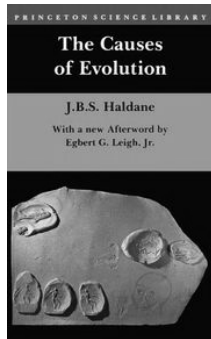
$$p^2 + 2pq + q^2$$

$$q^2 = .16; q = .4; p = .6$$

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Causes of Evolution

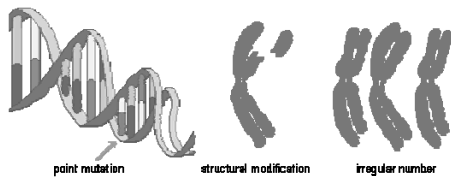
Non-random mating
Genetic drift
Mutation
Gene flow (migration)
Natural selection



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Mutation

point mutations
chromosomal
abnormalities



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Causes of Mutations

- ❖ DNA replication errors
- ❖ exposure to radiation, alcohol, lead, lithium, organic mercury, viruses, microorganisms
- ❖ teratogens: inhibitors, streptomycin, tetracycline, and vitamin A
- ❖ spontaneous irradiation, chemicals, temperature extremes

Mutation rate: 10^{-8} (1 in 100 million) per nucleotide per generation or 1 in 10,000-100,000 per sex cell

raw material for evolution

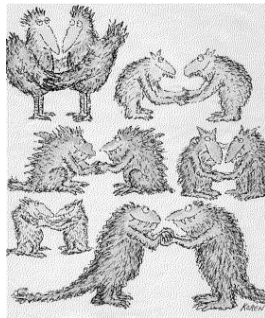
❖



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Non-Random mating

- ❖ non-random:
 - assortative mating (positive, negative)
- ❖ consanguineous marriage (inbreeding) e.g., Amish or Mennonites of New Foundland and Harrington Harbor in Quebec
- ❖ Since 1000 A.D., 2 raised to 39th power or 549,755,813,888 (5 1/2 trillion) ancestors
- ❖ incest prohibition
- ❖ increase homozygosity



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- ❖ genetic load: 3-5 lethal alleles
- ❖ E. g. rural Japanese: increased rates of congenital defects (higher rates of cleft palate and cardiac defects); pre-reproductive mortality rates increase; mental and physical growth are depressed

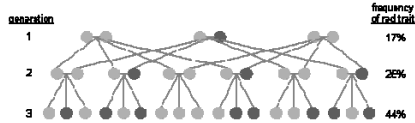


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Genetic Drift

Population Size and Sampling Error

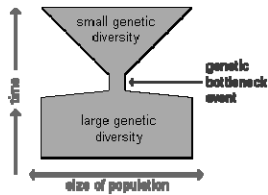
expected			chance deviation		
	A	a		A	a
A	AA	Aa	A	AA	Aa
a	Aa	aa	a <td>Aa</td> <td>aa</td>	Aa	aa



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Genetic bottleneck

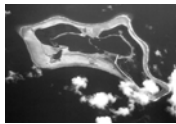
- when effective breeding population suddenly becomes reduced which might then drastically alter the genetic composition



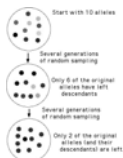
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Genetic Drift

E.g., Pingelap; typhoon 1775
 ♦ achromatopsia (1/60 or .016)



E.g., Alpine Villages (albinism, deaf mutes, blindness, mental deficiency)

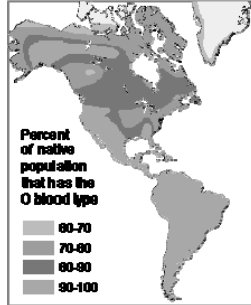


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Founder Effect or Founder Principle

E.g., Amish (200),
Hutterites (N=443),
Pitcairn Is.

Type O blood in South
& Central American
Indians



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Gene Flow

Hybridization and intermarriage – special kinds of gene flow
e.g., miscegenation; I^A and I^B alleles across Eurasia

admixture (e.g., American Black and American Whites)

Duffy blood group (Fy)

- ❖ Africans, Fy^o (a-b-) -- almost 100%
- ❖ Whites, Fy^o (a-b-) is rare -- 0%
- ❖ Fy^a = 43% in Whites and 0% in Africans

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Natural Selection

- ❖ selection for or against one of the following:
- ❖ either homozygote (AA or aa but not both)
- ❖ both homozygotes (AA and aa)
- ❖ either homozygote and the heterozygote (AA and Aa or aa and Aa)
- ❖ the heterozygote (Aa)
- ❖ all alleles (AA, Aa, and aa)



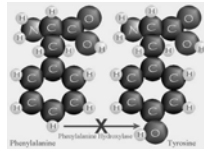
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Differential Mortality

death: eliminates individuals

- ❖ e.g. phenylketonuria (cannot metabolize phenylalanine)

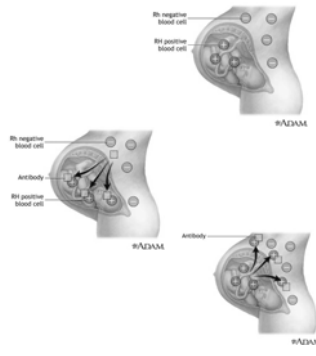
differential reproduction (or reproductive or survival selection)



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Rh incompatibility

- ❖ e.g. when mother is Rh- and offspring is Rh+: hemolytic disease of the newborn or *erythroblastosis fetalis*



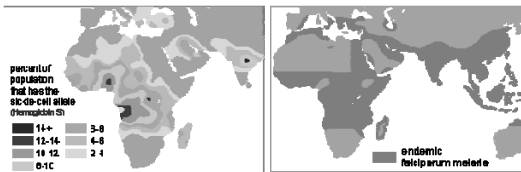
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Falciparum malaria & sickle cell hemoglobin

Hb^S Hb^S - sickle cell anemia

Hb^A Hb^A - normal individuals (die of malaria)

Hb^A Hb^S - sickle cell trait (mild anemia)



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