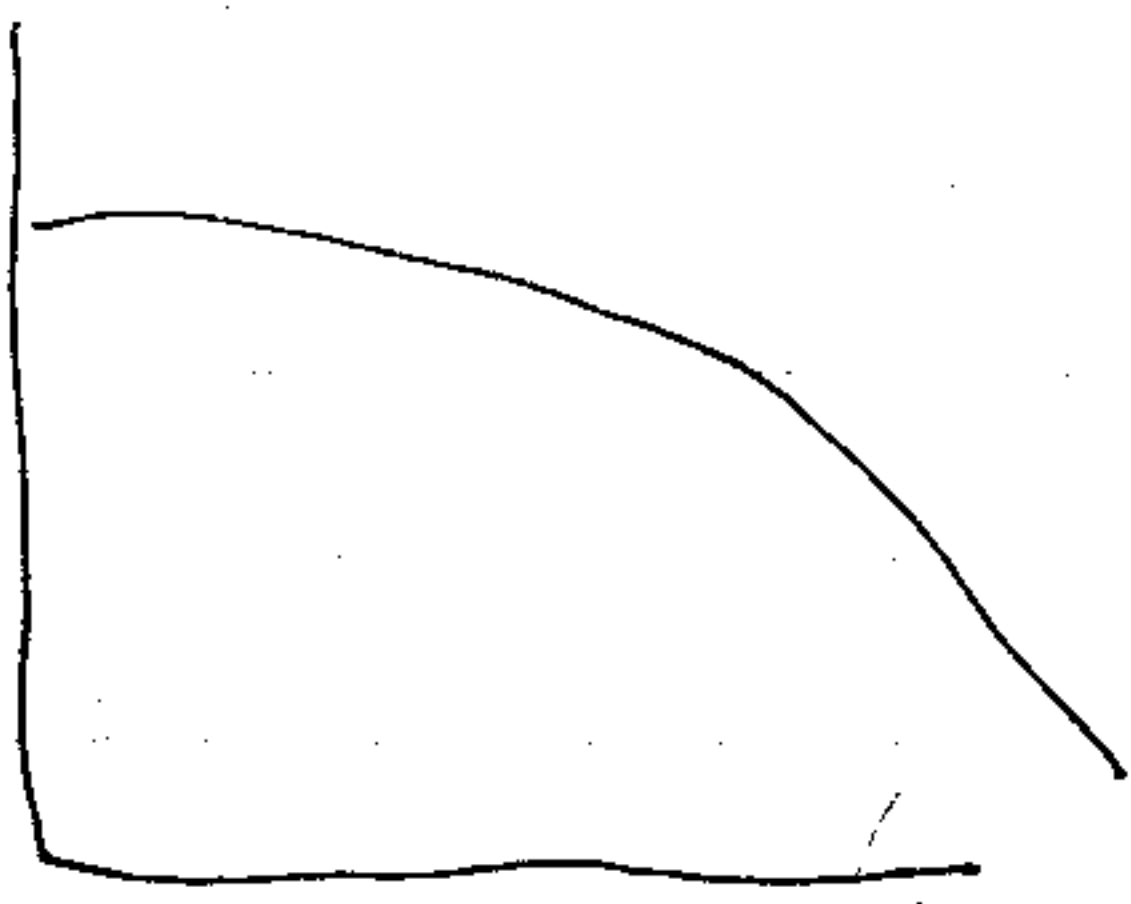


Fitness



additive scale

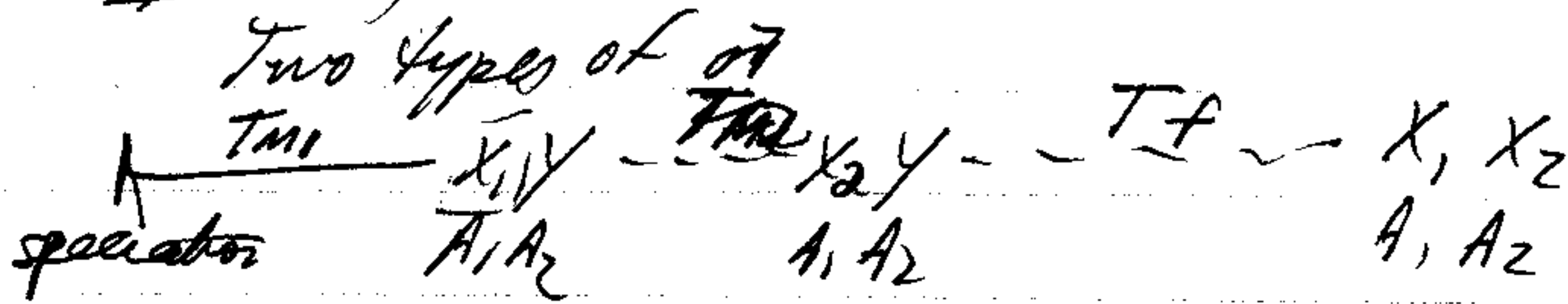
heterozygotes  
dominance parameter

$$b_{12} = h_1 b_{11} + h_2 b_{22}$$

X

The worse an allele is -- the more recessive it is.  
(generally the case in *Drosophila*)

Asymmetry



— don't always become inviable at same time of genetic separation

over time of speciation first one  $\sigma^7$  then other  $\sigma^7$  then  $\sigma^7$  becomes inviable

$X_1^* Y$   
 $A_1, A_2$  dead

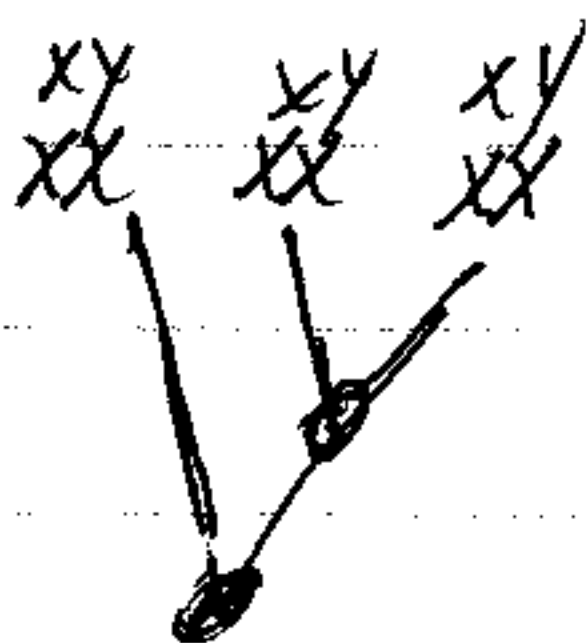
— the ratios of  $T_{m1}/T_{m2}$

$X_1^* X_2$   
 $A_1, A_2$

... w/ larger separation of  $T_{m1}$  species the closer  $T_{m1} + T_{m2}$  should be to each other

contingency recessive

if  $A_1, A_2$  then  $X_1^*$  is recessive lethal



— see if  $X_1^* Y$   
 $A_1, A_2$

SYNTHETIC LETHAL

M. Turrelli - Haldane's Rule

Haldane's Rule - heterogametic sex is inviable/sterile if one is

- Muller's

- heterogametic have only one allele for all X loci thus it is homozygous for recessive deleterious alleles

- Coyne (Coin?) - tested

- Flies

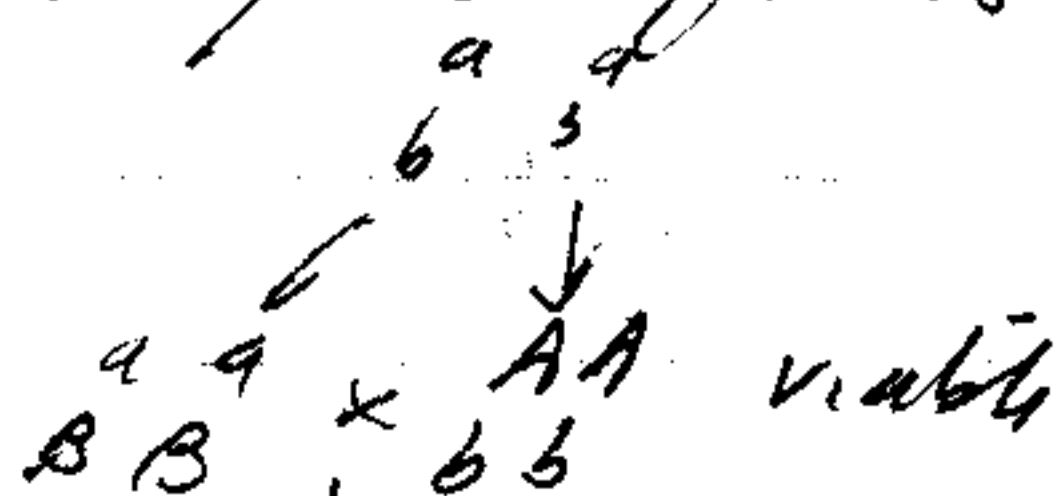
<u>hybrid</u>	{	$X_1 Y$	$X_1 X_2$	<u>hybrid</u>
		$A_1 A_2$	$A_1 A_2$	
		sterile	fertile	

- so if due to homo-recessive a ♀ w/ ~~AA~~ X<sub>0</sub> from same species should be sterile

- but not great test bec. ♂ + ♀ have diff. loci determining fertility

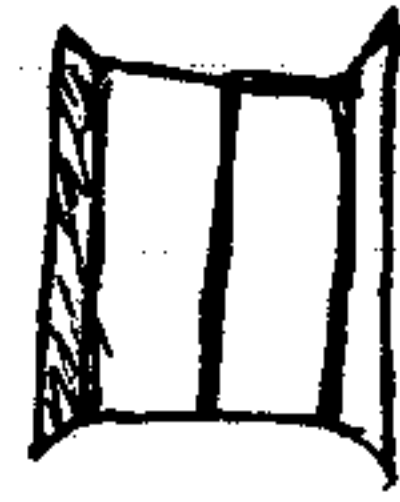
- so should do this test w/ viability - bec. some loci affect viability

- Why are hybrids inviable?



$\begin{matrix} A & A \\ B & B \end{matrix}$  - inviable because A & B don't interact well

⑦ Gordon Fox - Freq. Data & selection



Frequencies  $\xrightarrow[\text{to rates}]{\text{difficult here}}$  Process

Difficulties

- ① don't know about ~~the~~ equil.
- ② don't know about history of selection

\*- but says if know phenotypic affect of genotype can do OK -- but assumes this has been constant

⑧ Mark Bote - Sampling & Inference when a gene is under selection ...

Goal - char. type & intensity of selection if any

HLA

- fewer than expected homozygotes (but not many)
- neutrality can frequently be rejected by Watterson test
- assumed to be under balancing selection
- so need NULL SELECTION model

Watterson model

# High Salmon - UCB

Finds features that distinguish one sequence from another

seq	1	2	3	4	5	6
check seq	A	A	C	B	A	A
others	A	A	B	A	A	A
	A	A	B	A	A	B
	B	A	B	B	A	B
	C	A	A	A	B	B
	B	A	A	A	B	A

$\frac{245}{\text{---}} \leftarrow 4 \leftarrow$ 
 $\frac{245}{\text{---}} \leftarrow 5 \leftarrow$

Can be used for degeneracy

ART WEISS - G. C. Irvine

Genetics of Defense in Depth

- the fitness curve for a resistance allele depends on the tolerance
- Goldenrod & beetle

Susan Edmonds - UCSD - Urchin Genetic Structure



- sampling diff. ages important
- multiple markers important

*S. franciscanus*  
*S. purpuraceus*



- in some pops. adults & recruits have <sup>slightly</sup> diff. isozyme pattern
- allozyme - significant heterozygosity
- and - low hetero

Karen Marchetti - why do  $\sigma^7$  have multiple song chers

Warblers  $\sigma^7$

- bright patches

- song

why multiple?

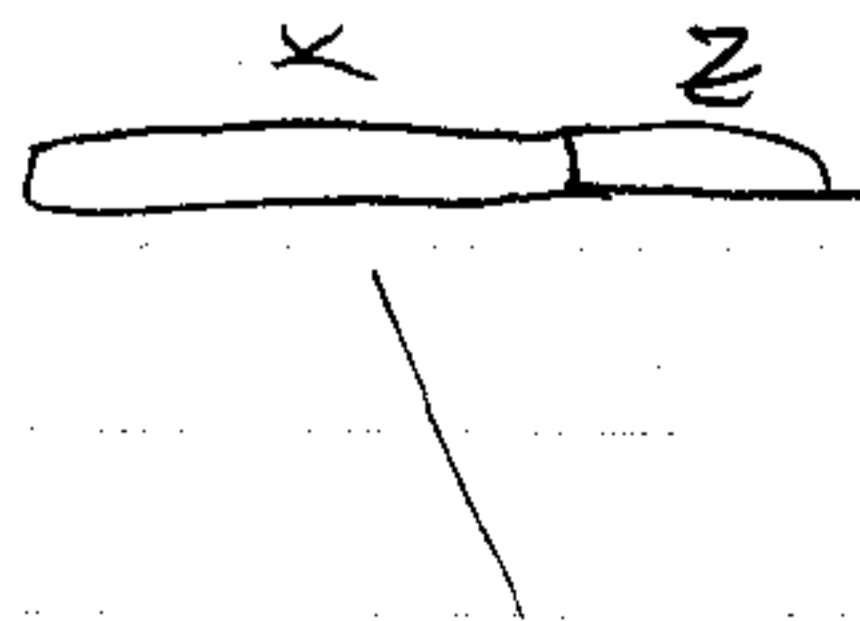
- ① ♀ choice may differ from  $\sigma^7$  compet.
- ② ♀ may choose many.
- ③ diff. chers may reflect diff quality.
- ④ variable ♀ choices.
- ⑤ diff signal context.

measurements

manipulation of color patch size

IS COLOR  
PATCH SIZE  
CONSTANT?

WHAT ABOUT  
♀ SEEING  
THROUGH  
PAINT?





Grant Pogo - Comparing Protein & DNA Polymorphism

Comparing DNA & protein

① intra-locus

- e.g. Adh (MacDonald & Kreitman)

② inter-locus

- e.g. Karl & Avise (1992)

- argue that diff. in pattern of allozymes vs. DNA is due to selection

Uniformity of process

Drift

Equal ??

Mutation?

Migrations

Equal

Inbreeding

Equal

Selection

Variable

Mutation?

Markers

- transcribed (use cDNA RFLPs)

- mutation rate  $10^{-6}$  -  $10^{-7}$

- BUT THIS IS BIASED TOWARDS ABUNDANT ALLELES

Heterozygotes

Agents

Overdom.

Multi-locus dom.

mut alleles

Impurity

Markers

Assoc. overdom.

Inbreeding

Somatic aneuploids

Is gene flow important

① gene flow should affect all loci equally

② so polymorphism should be same at all loci

HOW SELECT FOR RARE ALLELES

MUTATION?



# CALPEG 1994

Trevor Price

see Felsenstein }  
1985 }  
- w/ biological correlations it is generally assumed that phylogeny must be taken into account to determine significance of association

- unless of course selection is very effective and quick - that is all things are optimized

- another problem is if another character affects the traits - <sup>only one of</sup> only

- Felsenstein method corrects for phylogeny

BUT THESE ARE STILL VERY CLOSE RELATIVES }  
- but Trevor says nothing affects only one variable (e.g. brain & body size both affected by same selection)

- so CONTRASTS (FELSENSTEIN) ASSUMES PHYLOGENY IS TOO IMPORTANT