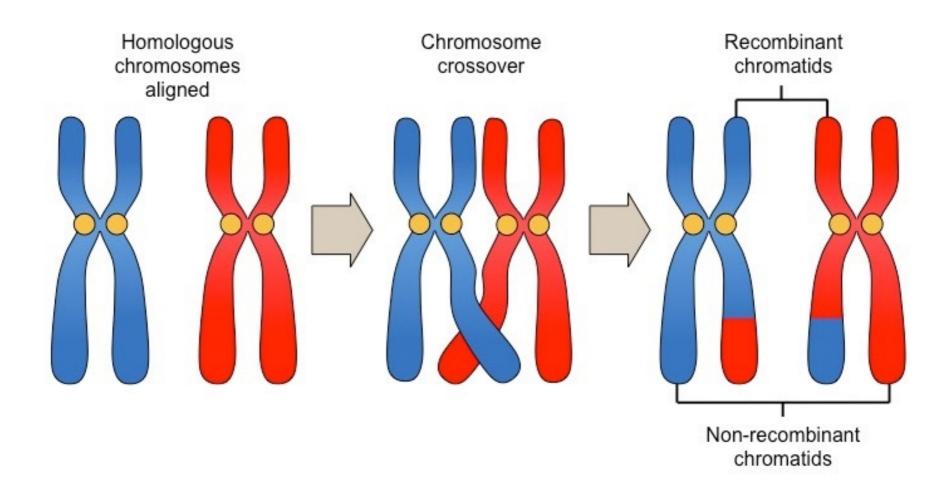
Interactions between several loci, Epistasis, Super Genes, Pleiotropy

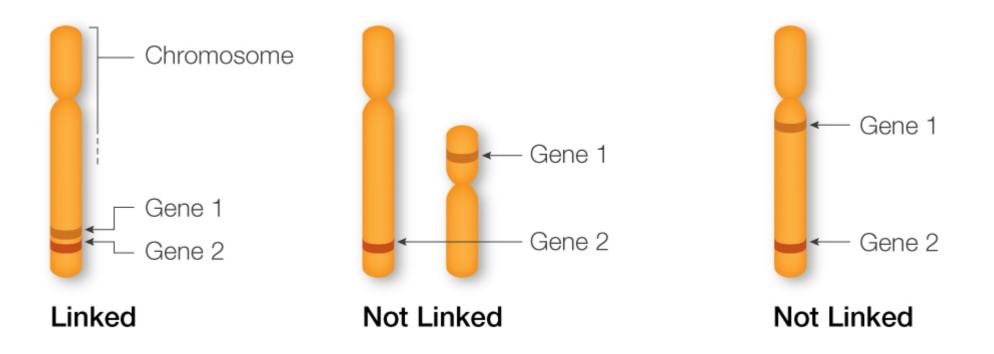
Virginie Courtier-Orgogozo Institut Jacques Monod, Paris

Genetic Linkage

Crossing overs



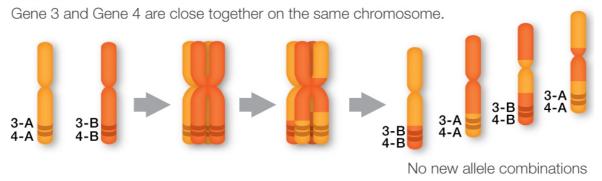
About one recombination event per chromosome arm



https://learn.genetics.utah.edu/content/pigeons/geneticlinkage/

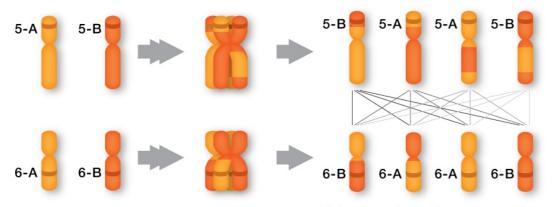
Not Linked Gene 1 and Gene 2 are far apart on the same chromosome. 1-A 1-B 1-A 1-B 1-A 1-A 2-B 2-A 2-A 2-B 2-A 2-A New allele combinations

Linked



Not Linked

Gene 5 and Gene 6 are on separate chromosomes.



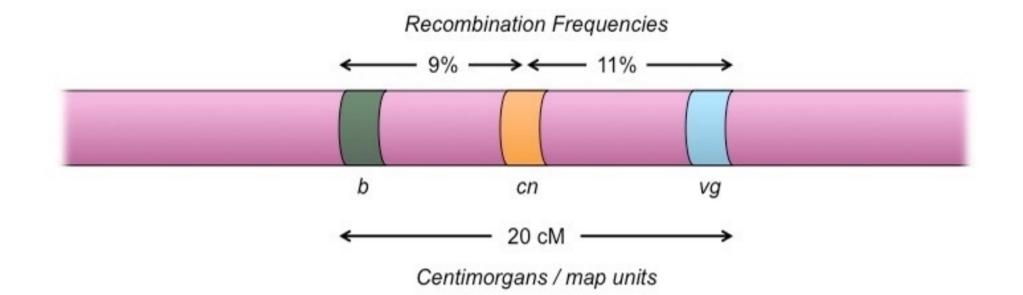
Alleles (on whole chromosomes) can be distributed to gametes in any combination.

One "centiMorgan"

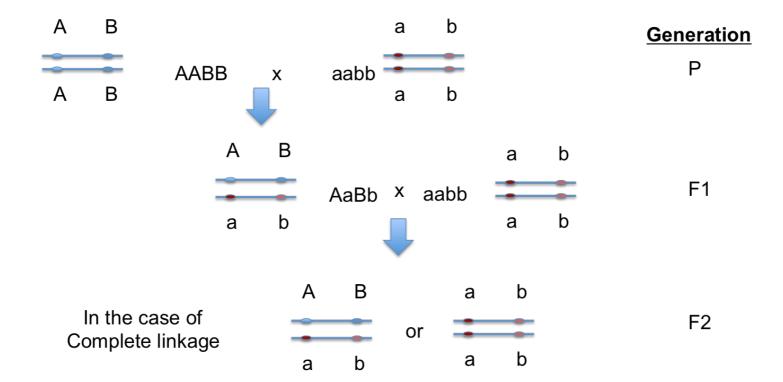
= genetic distance that produces a recombination frequency of 1%

Genetic distance (in cM)

= <u>(# Recombinant gametes) X 100</u> Total gametes



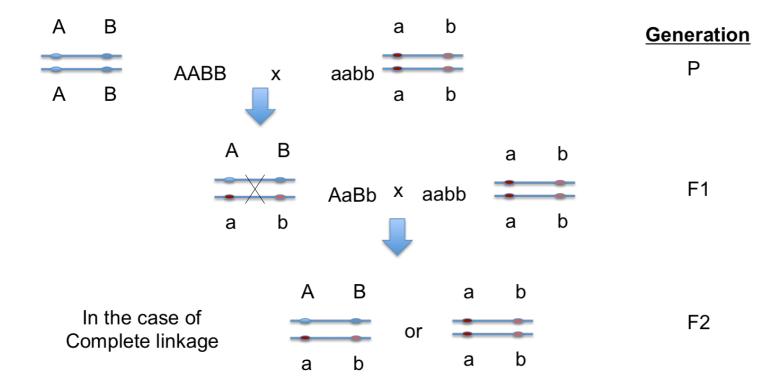
Measure of genetic linkage



Complete Linkage

50% AaBb 50% aabb

Measure of genetic linkage



Complete Linkage

50% AaBb 50% aabb

Genetic Linkage

40% AaBb 10% Aabb 10% aaBb 40% aabb

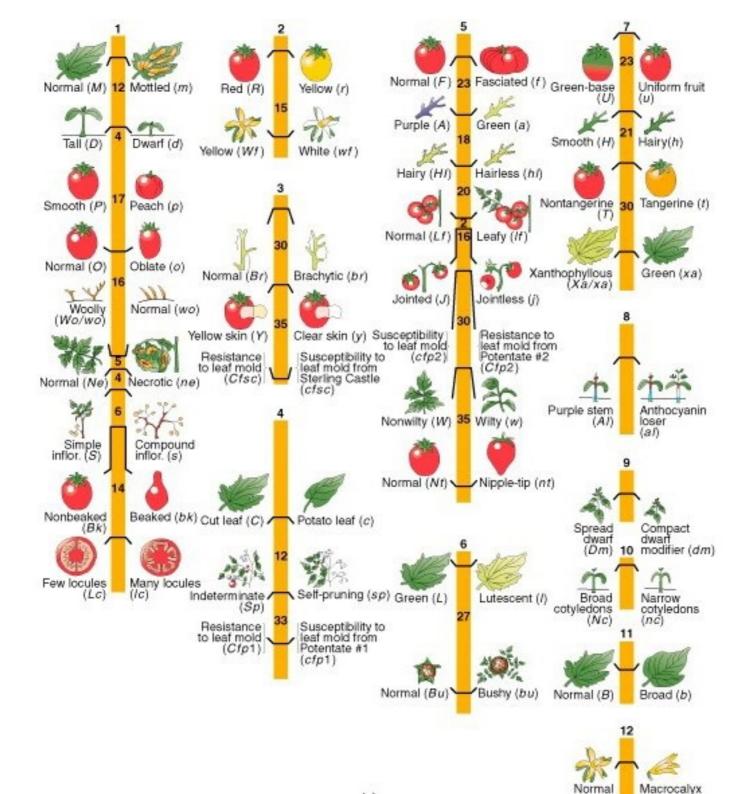
20% of recombinants so 20cM

Measure of genetic linkage

If y % recombinant gametes and y < 50% => y cM apart

Due to double cross-overs and cross-over interference, genetic distances need corrections when long and are not fully additive

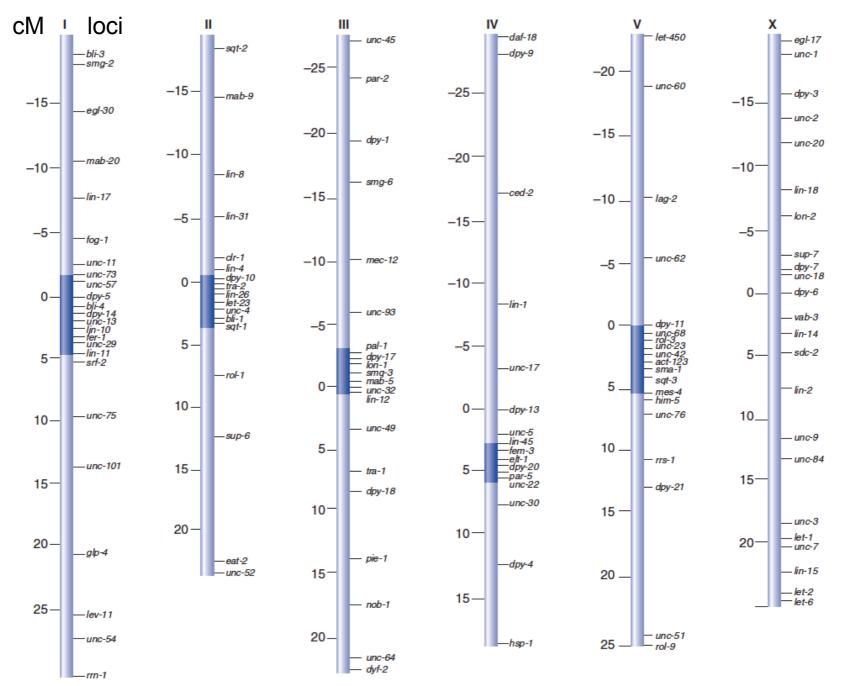
If the linkage group is longer than 50 cM, mutations at the two extremities are operationally unlinked



Genetic map

in units of recombination

1 centiMorgan (cM) = 1% recombinants



Genetic Markers

Mark the region of interest through genetic linkage

Are not causal (or only rarely) for variation in the phenotype of interest

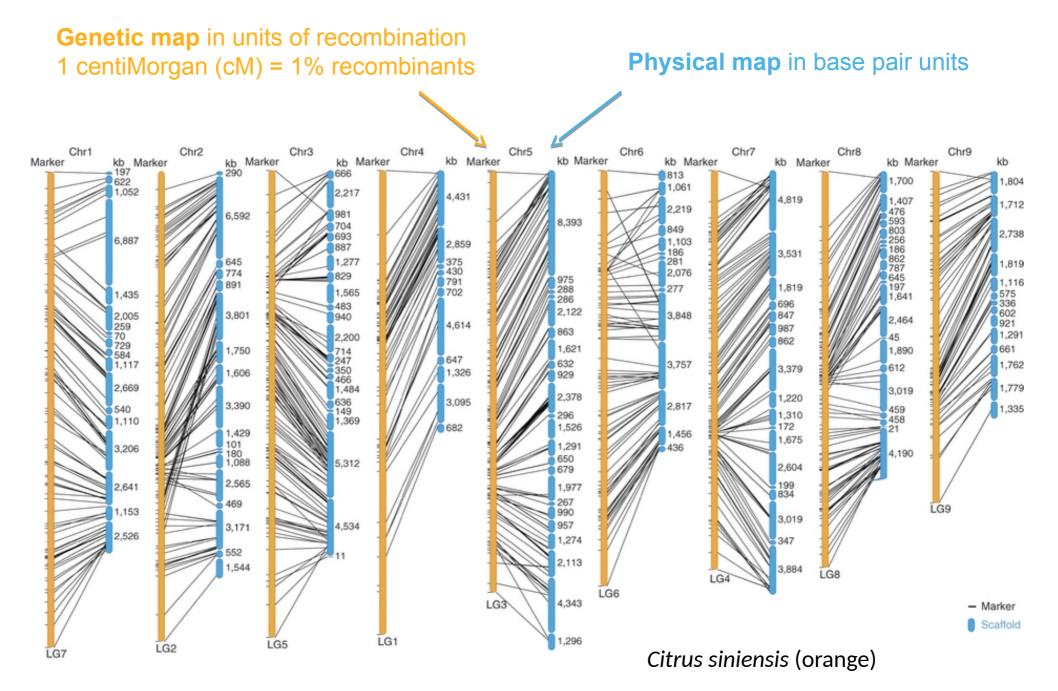


Detected:

- through their phenotypic effect: white eyes, dumpy shape, GFP marker

- molecularly: PCR, sequencing transposon insertion, single-nucleotide polymorphism (SNP), indel

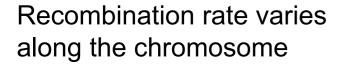
Alignment of genetic and physical maps



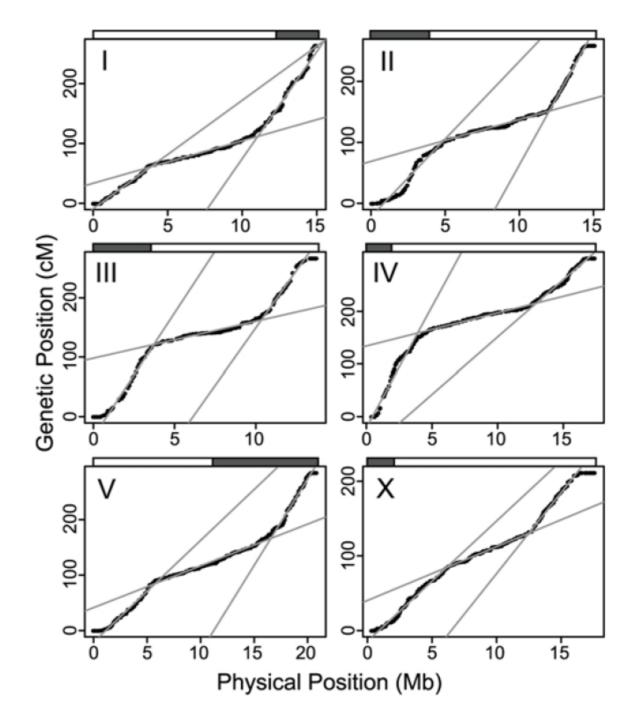
Alignment of genetic and physical maps

Marey map

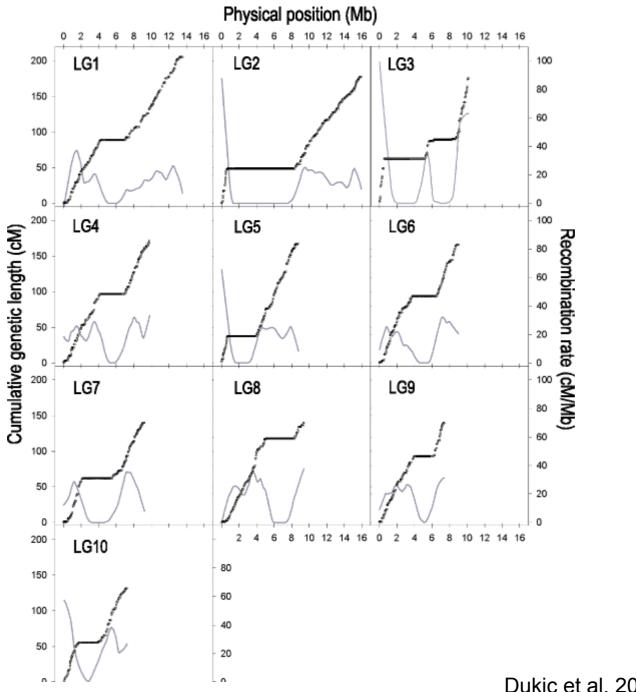
Genetic position was measured in centiMorgans based on a recombinant inbred advanced intercross line population, and not based on meiotic distances.



C. elegans Rockman & Kruglyak *PLoS Gen* 2009



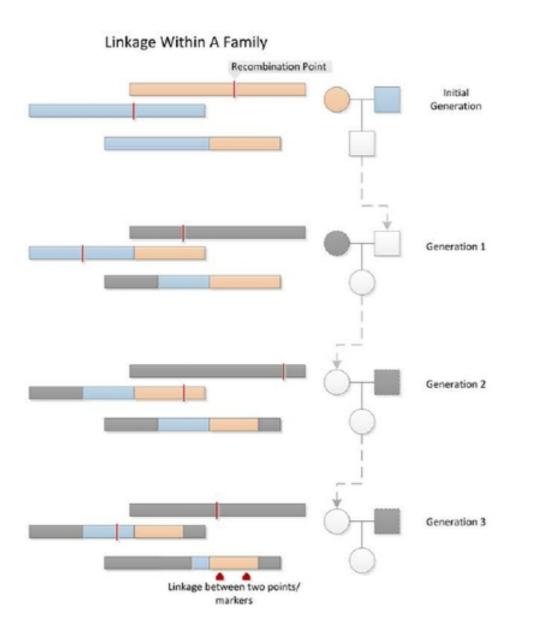
Marey maps in Daphnia



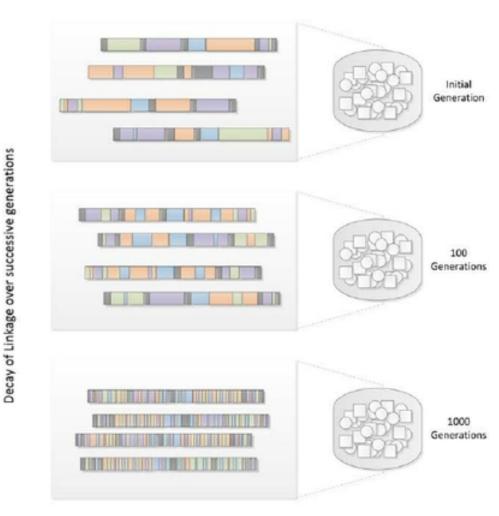
Dukic et al. 2016 BMC Genetics

Linkage disequilibrium (LD)

non-random association of alleles at different loci in a given population



Linkage Disequilibrium Within A Population



Population moves from Linkage Disequilibrium to Linkage Equilibrium over time

Gokcumen et al 2013 NE1 Locus (36 kb) chr22:37,600,063-37,636,026 37,600,000 37,640,000 37,560,000 37,580,000 37,620,000 37,660,000 37,680,000 CBX6 APOBEC3A CNVR8163.1 Deletion 37624055 3762863 nonNE1 Haplotype TTGTTTTATTTTAAAGTGAAAACTGTAA TGGGGCAAACAGCTTTCCT GGGGCAAACAGCTTTCCT Mbut NA10494 TTGTTTTATTTTAAAGTGAAAACTGTAA CEU NA12155 TTGTTTTATTTTAAAGTGAAAACTGTAA GGGGCAAACAGCTTTCCT CEU NA12008 TTGTTTTATTTTAAAGTGAAAACTGTA GGGGCAAACAGCTTT Surui NA10970 TTGTTTTATTTTAAAGTGAAAACTGTJ GGGGCAAACAGCTT Surui na10972 TTGTTTTATTTTAAAGTGAAAACTGTAA GGGGCAAACAGCTT Mayan NA10976 TTGTTTTATTTTAAAGTGAAAACTGT/ GGGGCAAACAGCTTTC Mayan NA10975 TTGTTTTATTTTAAAGTGAAAACTGTAA GGGGCAAACAGCTTT Pima NA14313 TTGTTTTATTTTAAAGTGAAAACTGTAA GGGGCAAACAGCTTTCCTT Pima NA14308 TTGTTTTATTTTAAAGTGAAAACTGTAA TGGGGCAAACAGCTTTCCTTAGC ¹36-kb NE1 haplogroup contains a 4.6-kb deletion in perfect linkage disequilibrium with 12 SNP aligns with Neandertal haplotype

The Linkage Disequilibrium (LD) block was determined using SNP data of the CEU population from 1000 human Genomes

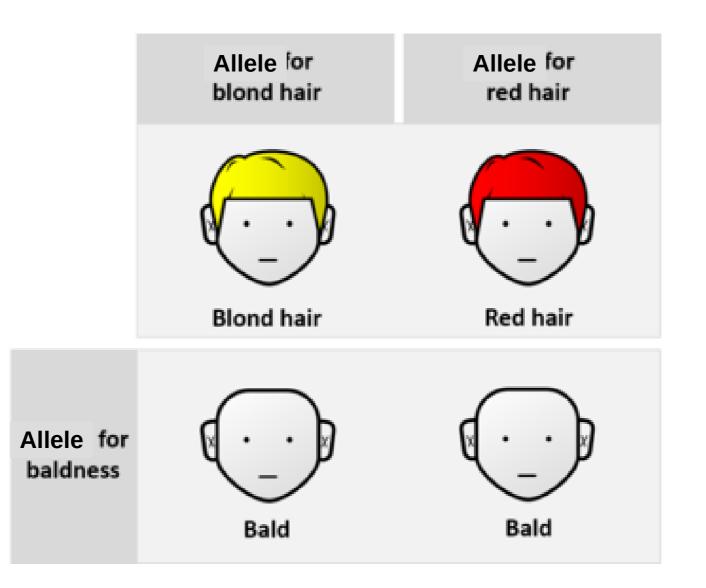
Variation in Linkage disequilibrium (LD)

LD is a function of age of alleles, outcrossing and recombination rates

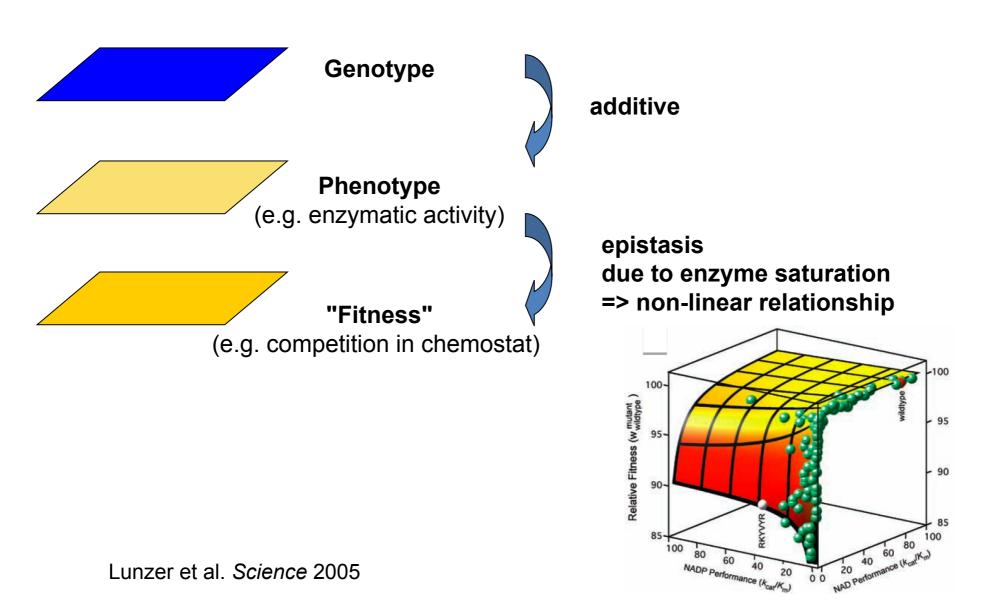
Depends on organism and genome region short-range = 100 bp *D. melanogaster*, *Caenorhabditis remanei* medium-range = a few kb: *Homo sapiens*, *Arabidopsis thaliana* long-range = Mb: *Caenorhabditis elegans*

Epistasis

= Non-additive interaction of alleles at different loci for a given phenotype



Additivity at one phenotypic level does not imply additivity at another level



Various meanings for Epistasis

Laboratory genetics, with null alleles

m1 is epistatic to *m2* if *m1 m2* displays the M1 phenotype => genetic pathway

Quantitative / evolutionary genetics

"epistasis" used for "gene interaction"

= non-additive effect for any combination (heterozygote, homozygote) non-additive mapping of genotype space to phenotype space

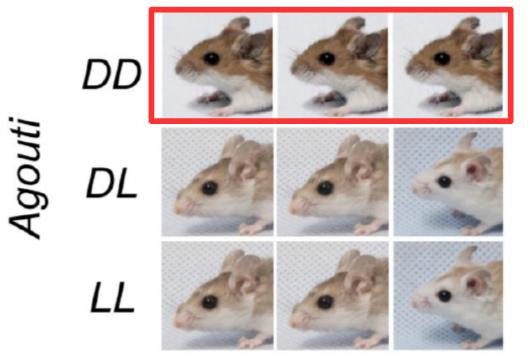
=> confusion between lab geneticists and evolutionary geneticists

Meaning of "epistasis" depends on the scientific context!

Agouti (D, L) and Mc1R (D,L)

Natural alleles 3 phenotypes

DD DL LL



Agouti^D is epistatic over Mc1R alleles

Philipps 2008

Agouti (A, a) and Mc1R (E,e)

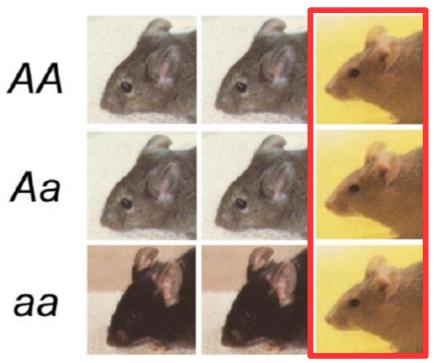
Laboratory mutants 3 phenotypes







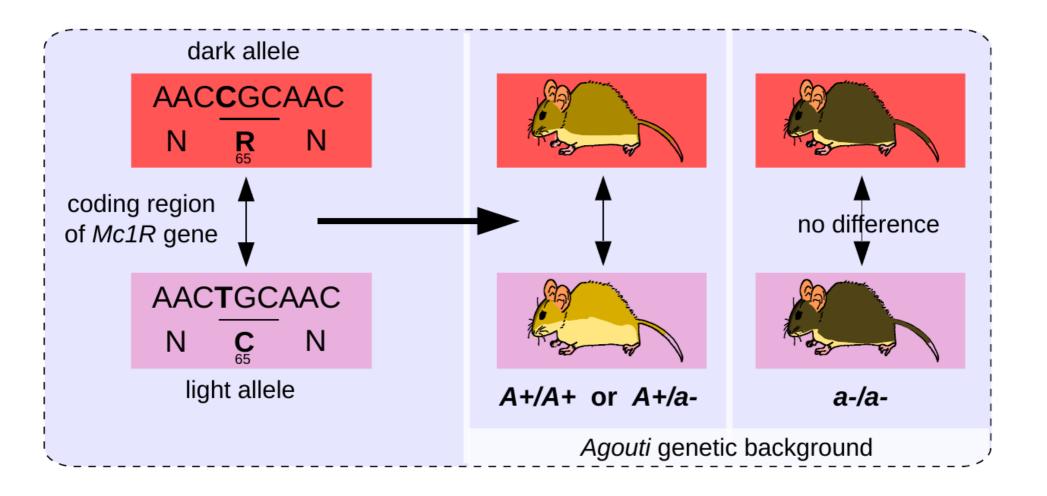
EE Ee ee



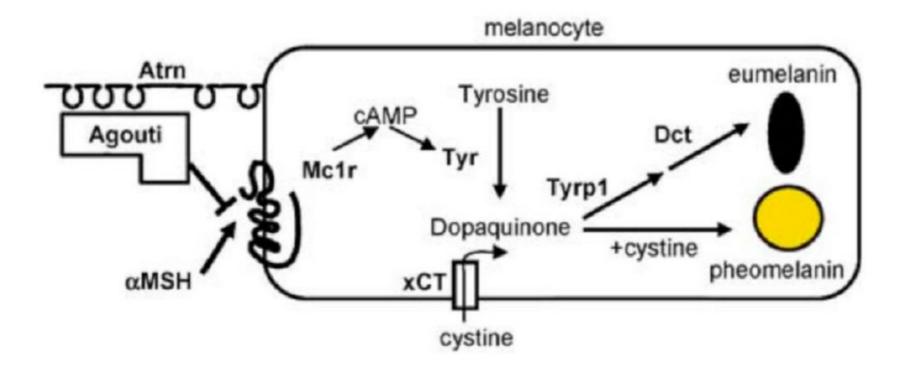
Agouti

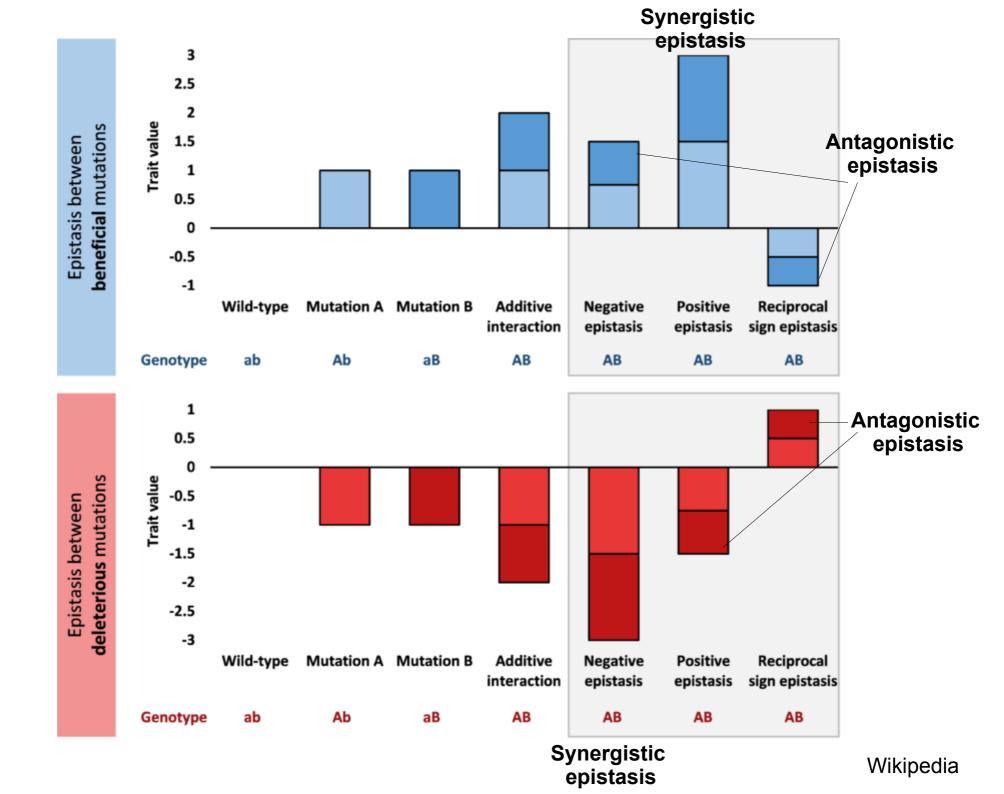
Mc1R^e is epistatic over Agouti alleles

Philipps 2008



Orgogozo, Morizot, Martin 2015



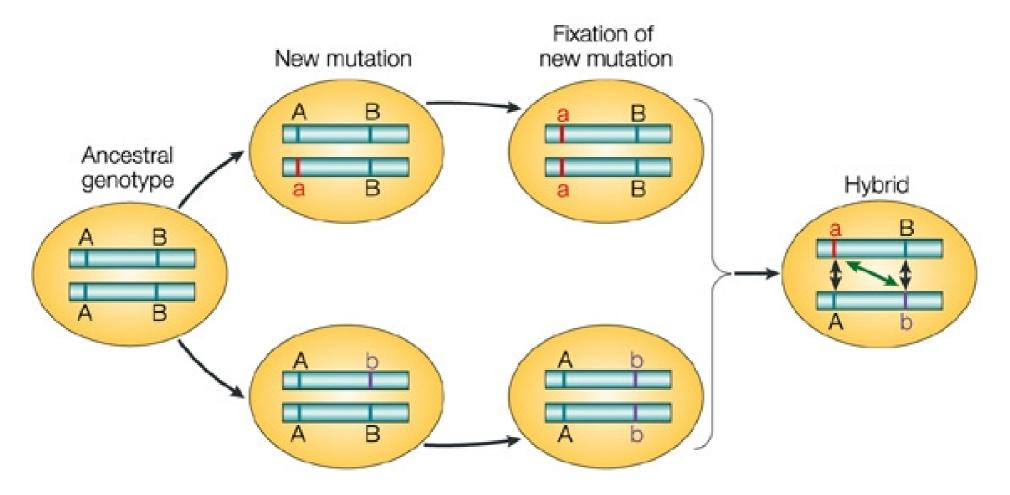


"Synergistic epistasis": the effects of both alleles reinforce each other (more than the sum of their individual effects); extreme case: synthetic phenotype (new phenotype)

"Antagonistic epistasis": the effects of the two alleles partially compensate (less than the sum of effects of *a2, b2*)

"**Positive or negative epistasis**": the <u>phenotypic value</u> is either increased or decreased relative to additivity

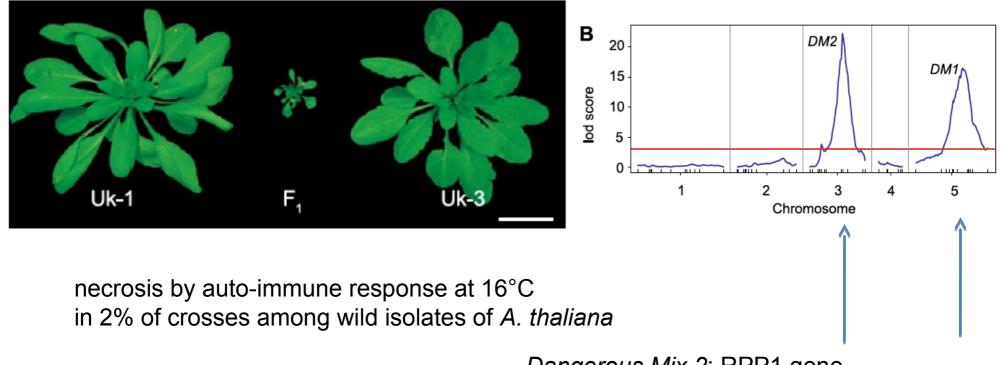
Dobzhansky-Muller model of hybrid incompatibility A special case of epistasis



possible mechanism of speciation

Nature Reviews | Genetics

Hybrid incompatibility in A. thaliana



Dangerous Mix 2: RPP1 gene resistance against oomycete

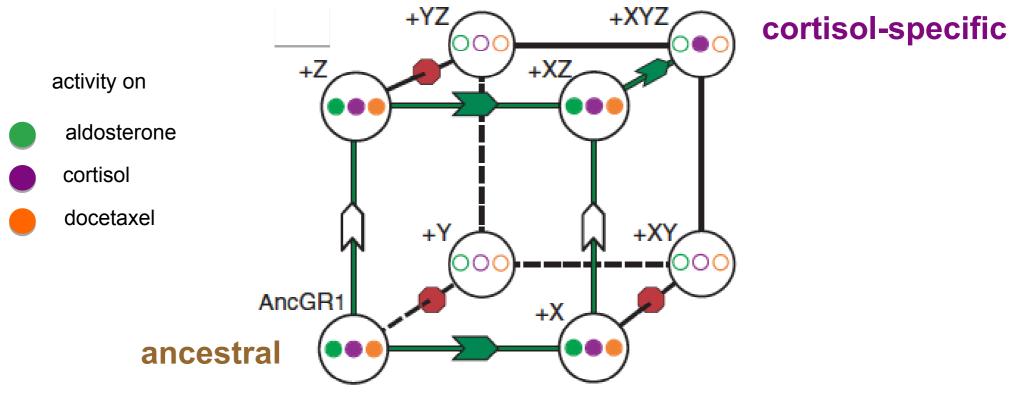
Dangerous Mix 1: member of a large family of pathogen resistance gene NB-LRR

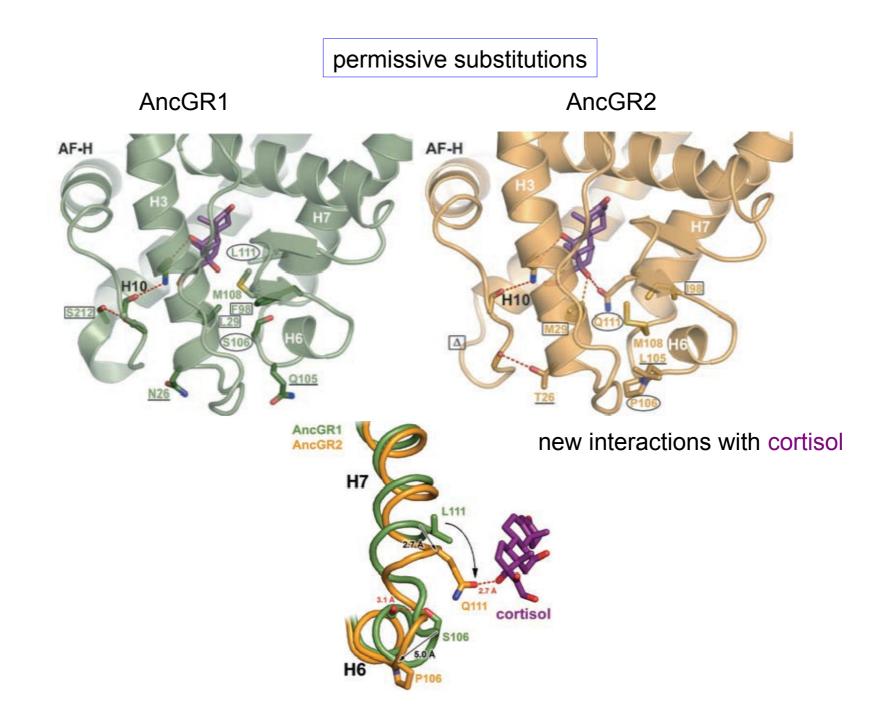
Bomblies et al. PLoS Biology 2007, Chae et al. Cell 2015

Intramolecular epistasis

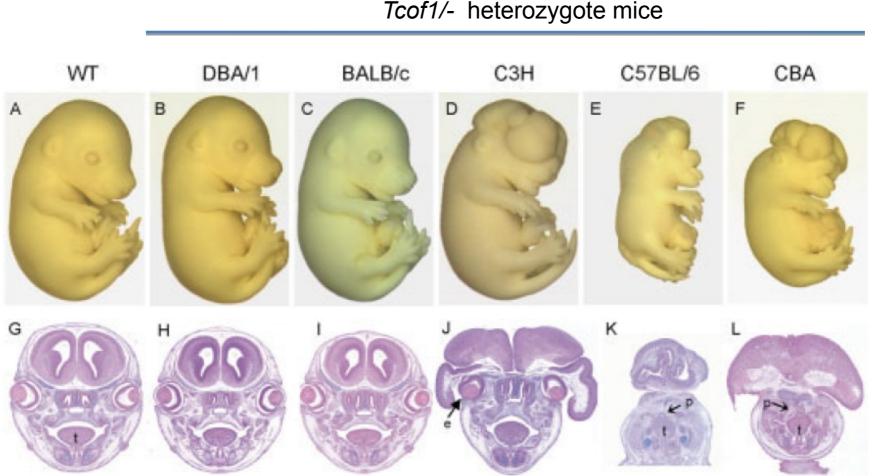
Reconstruction of ancestral protein sequence from phylogenetic analysis of extant family in databases

Vertebrate corticoid receptor family





Expressivity of one mutation varies with wild genetic gackground



 $f1/_{-}$ heterozyante mice

Dixon & Dixon Dev Dyn 2004

Different kinds of GxG interactions

- **G** x **G** between 2 laboratory mutations
- m1 m2

- G x G between 2 natural alleles
- a1/a2 b1/b2
- G1 x G2 one mutation ^m ^m ^m ^m ^{one mutation} in different wild genetic backgrounds "cryptic" variation

$G \times G \times G >2$ loci

a1/a2 b1/b2 c1/c2

Gerke et al. 2010

Super genes

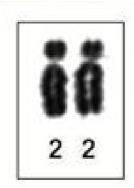
White throated sparrow

White morph





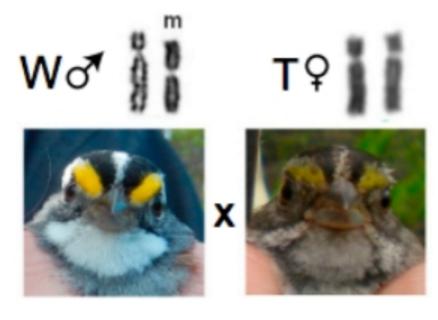
<image>



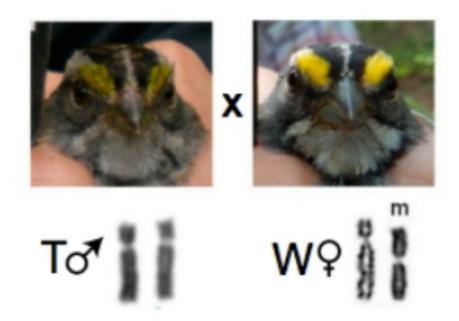


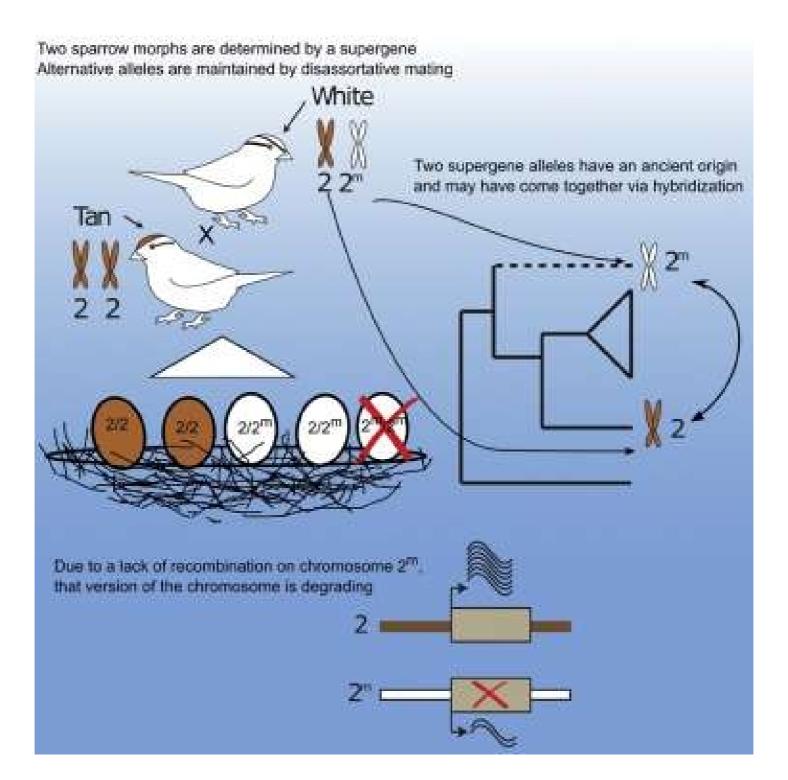
Tuttle et al. 2016 Curr Biol

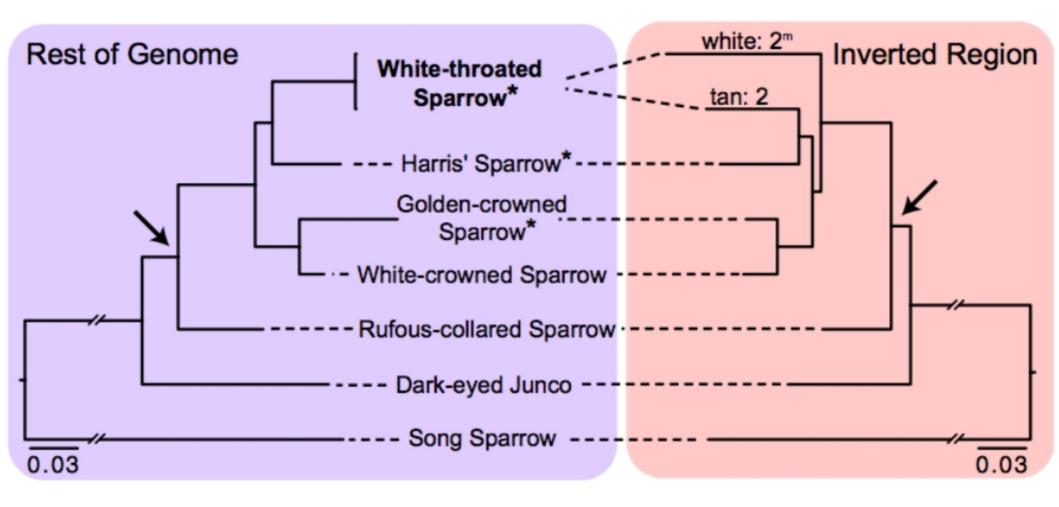
Disassortative mating



Never W male x W female Never T male x T female

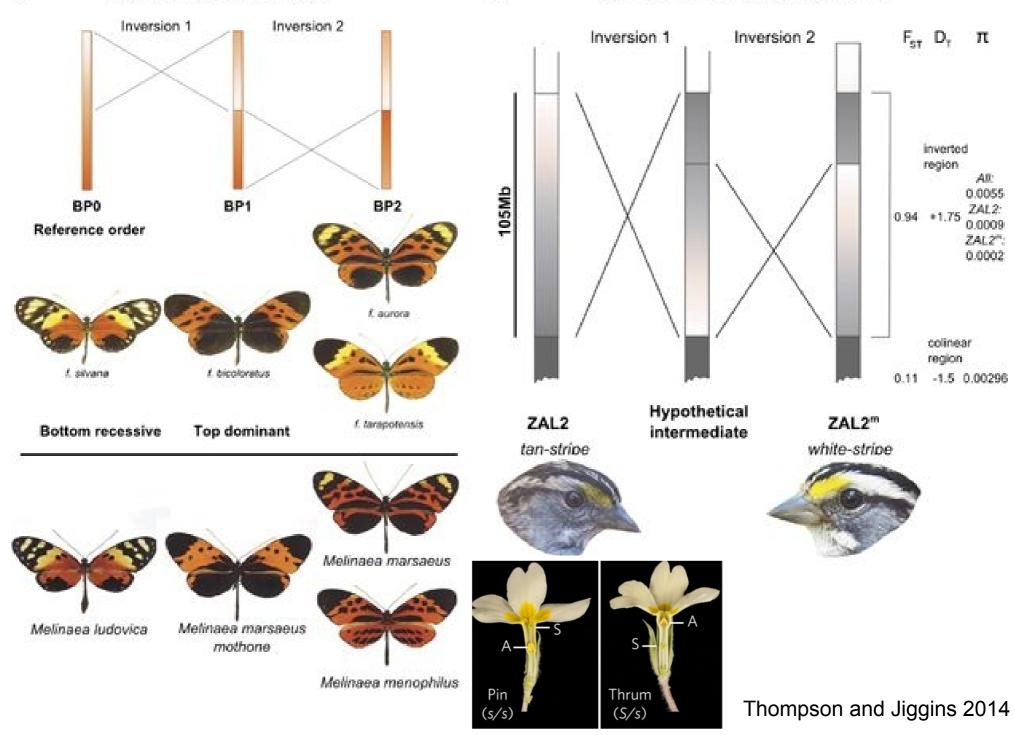






Heliconius numata P supergene

Zonotrichia albicollis chromosome 2



b

Pleiotropy

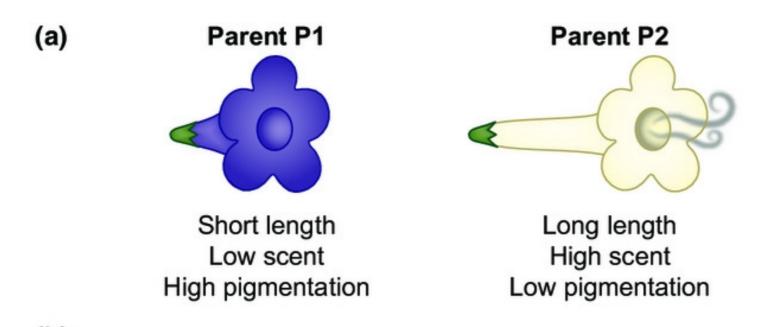
= when a genetic change affects several phenotypes

Various meanings for Pleiotropy

Pleiotropy of a gene (means pleiotropy of the *null* mutation)

Pleiotropy of a cis-regulatory region (means pleiotropy of the *deletion* of the region)

Pleiotropy of a mutation



(b)				
	Length	Scent	Pigmentation	
QTL1	1			No measured pleiotropy
QTL2	1	↓	\checkmark	Antagonistic pleiotropy
QTL3	1	1	♦	Adaptive pleiotropy

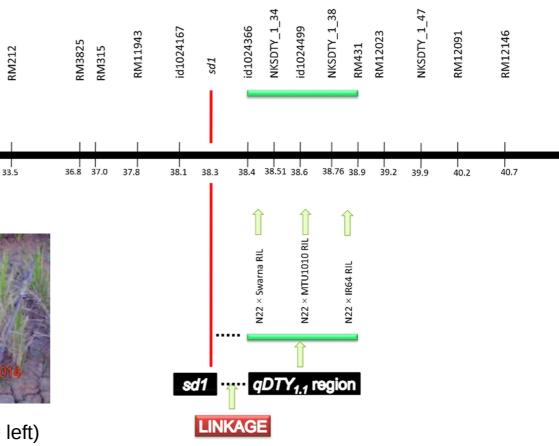
Modern rice varieties are sensitive to drought

sd1 locus (dwarf size) close to the *QDTY1.1* locus (grain yield under drought)



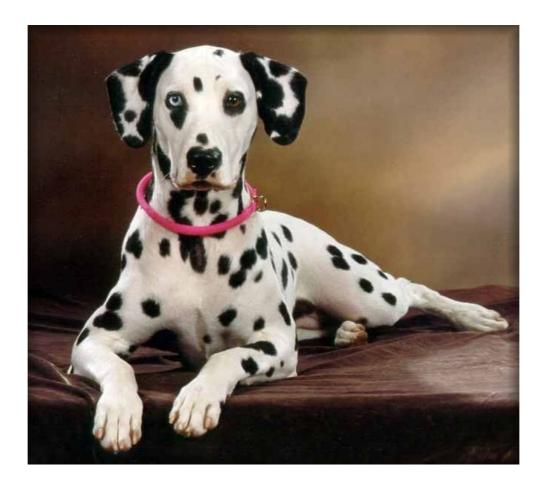
Lines with tolerant allele of qDTY1.1 QTL (on the left) remained green while those with sensitive allele (on the right) were severely affected under vegetative stage drought at IGKV, Raipur. Both the lines were of dwarf stature due to presence of sd1 allele.

Bhandari et al. 2019, Indian Society of Genetics and Plant Breeding



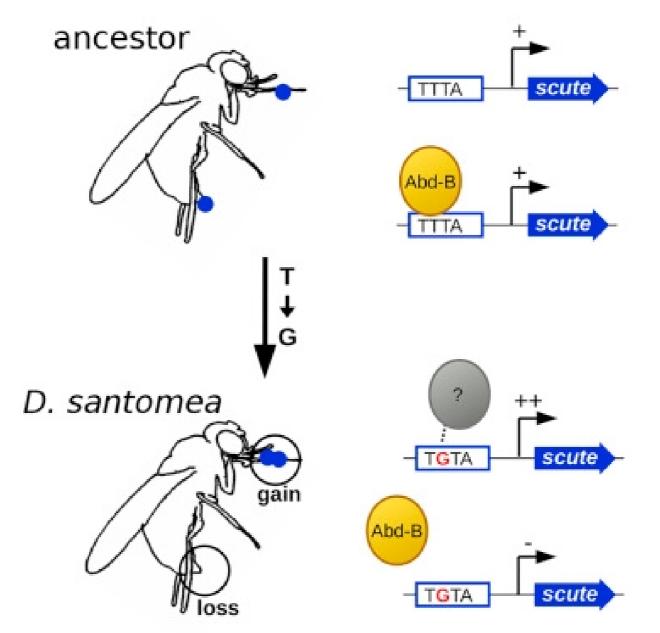
Vikram et al. 2015, Scientific reports

Dalmatian deafness



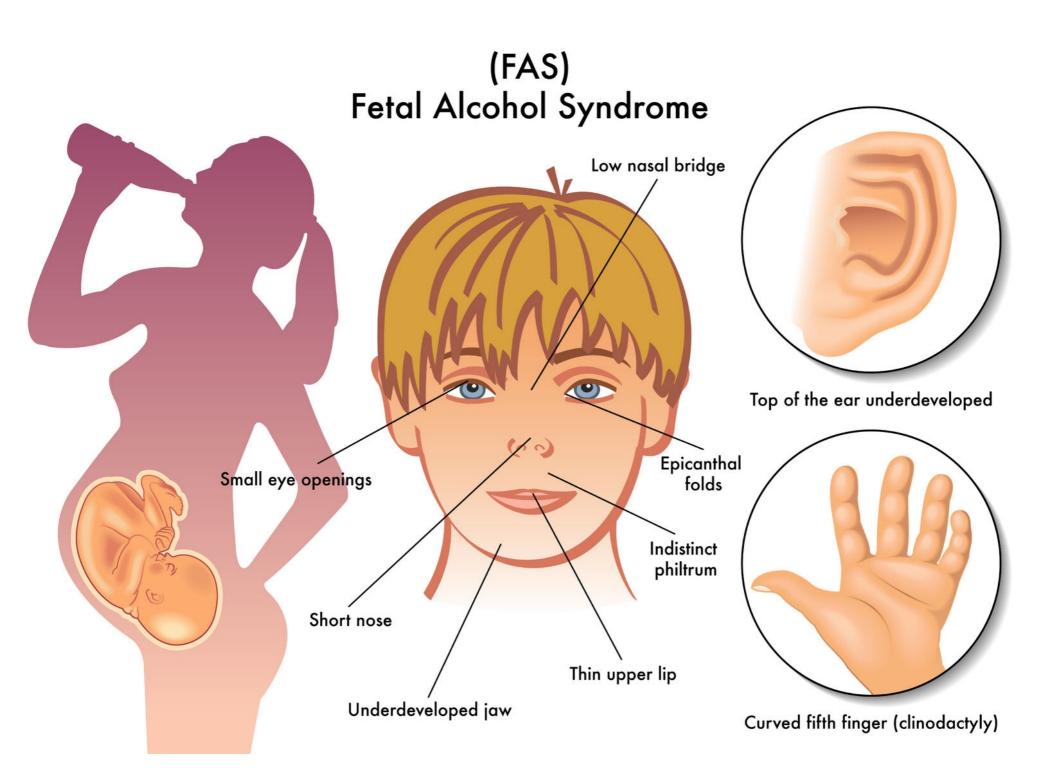
8% of all Dalmatians are bilaterally deaf and 22% are unilaterally deaf

A pleiotropic cis-regulatory mutation responsible for species difference



Nagy et al 2019

GxE



Causes of skin color differences

Genetic

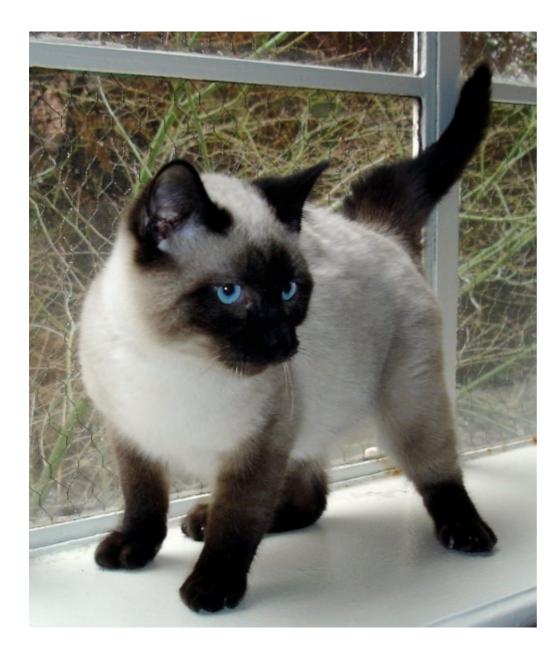
Environment





Phenotype = G + E + GxE

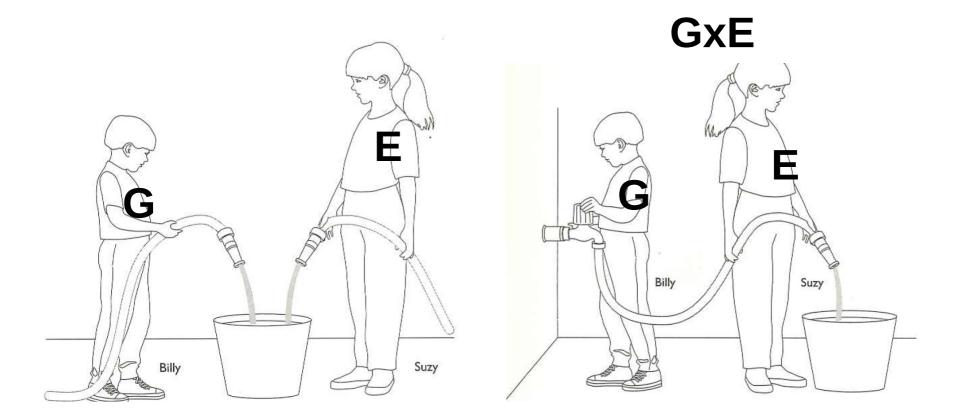
The Siamese cat An example of GxE



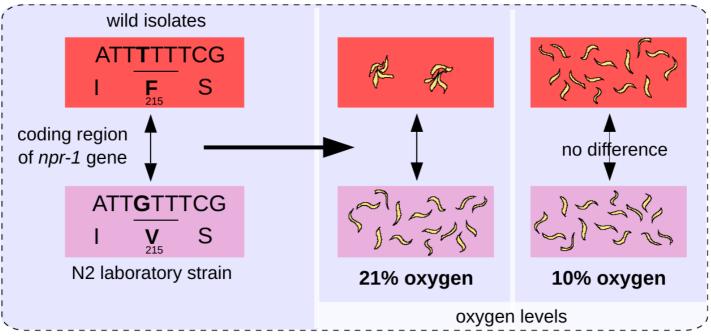


Mutation in *tyrosinase* Heat-sensitive enzyme No production of melanin in warm body parts

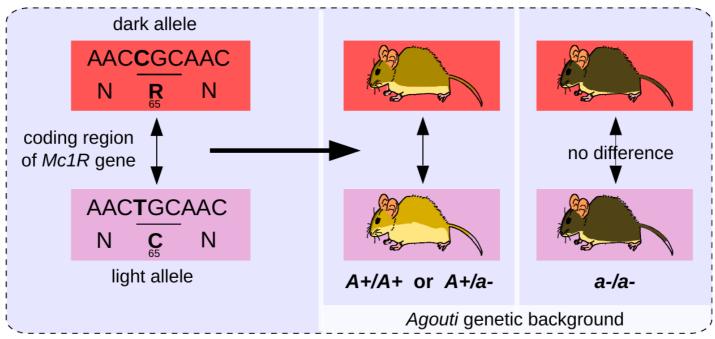
Contributions of the genotype (G) and the environment (E) to phenotypic variation



A GxE interaction

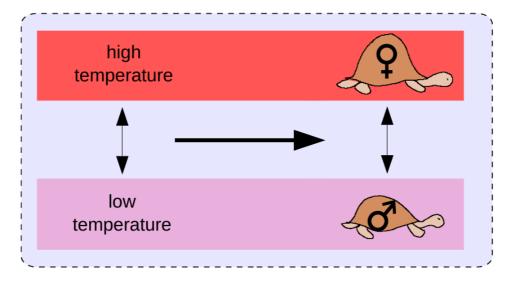


B GxG interaction

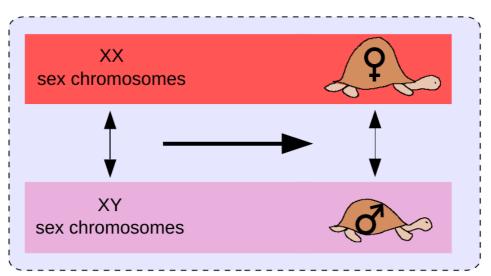


Comparing G and E effects

A enphe



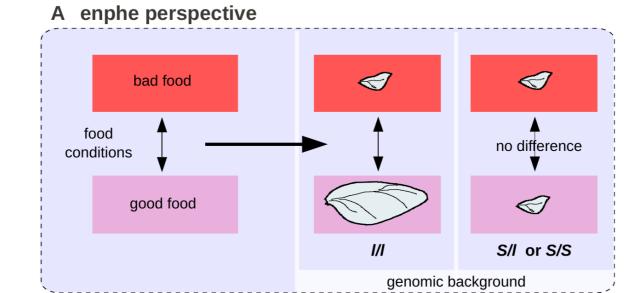
B gephe



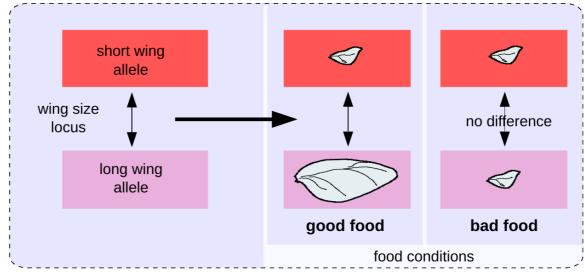
Intermingled G and E effects

Calathus melanocephalus

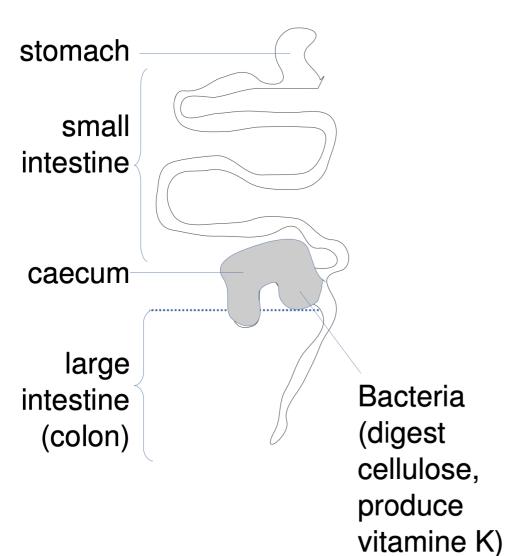




B gephe perspective



Mouse caecum development An other example of GxE





germfree

normal

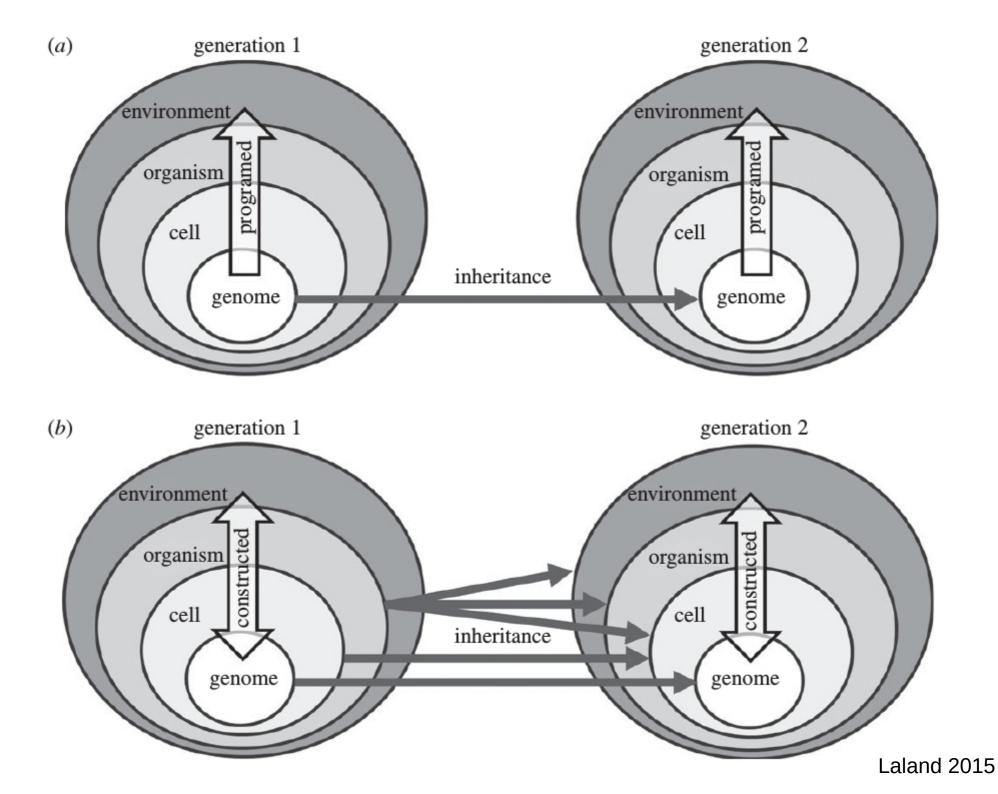
Causes of phenotypic differences? Heritable Non heritable





Phenotype = H + NH + HxNH

Like GxE but not always (Exceptions: DNA methylation, microbiome, langage, accent, culture, life style, parental care, maternal effet...)



Complexifications of the G-P map

Genetic Linkage

Epistasis

Supergene

Pleiotropy

GxE