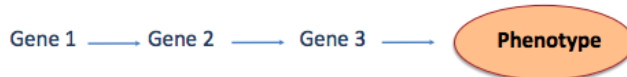


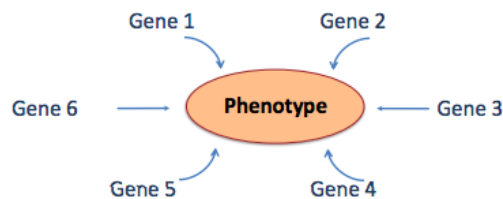
Lecture 5: gene interaction and complementation

How different genes contribute to a phenotype

- Linear or dependent functions=genetic interaction



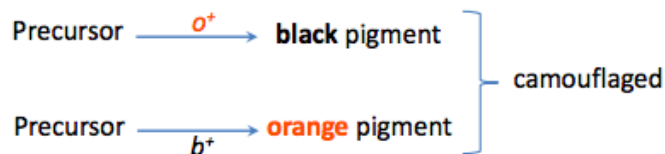
- Parallel or independent functions=synthetic interaction



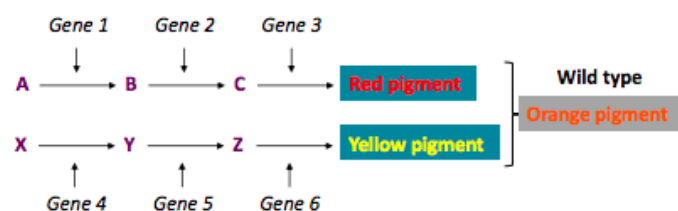
Variations in dihybrid cross

Example:

- a snake have 4 phenotype (WT camouflage, black mutant, orange mutant & albino mutant)
- Pure breeding black and orange snakes are crossed and all F1 progeny are WT(complement)
- F1 self crossed and in F2 all phenotype are found (9 WT : 3 black : 3 orange : 1 white)
- Camouflage is a dihybrid trait, there is no genetic interaction
 - > synthetic phenotype =created by allele of different genes in combination



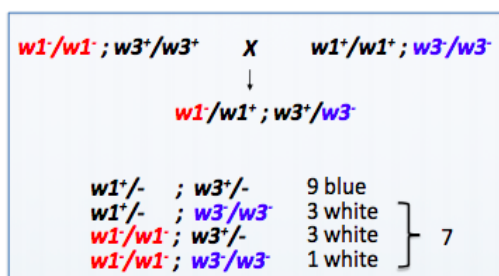
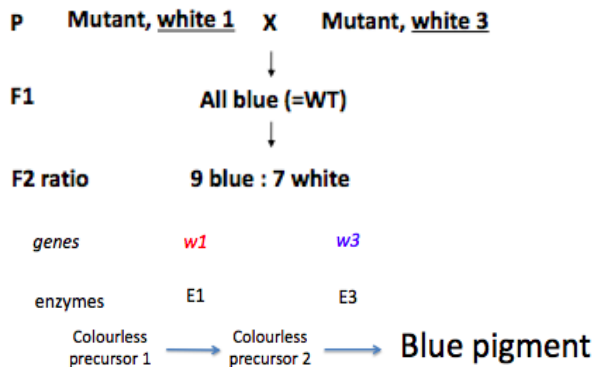
- There can be many precursors that leads to one phenotype
- Genes in the biochemical pathway affect the same phenotype



- Mutation in any of these genes alter pigmentation
- If at least 2 different genes from each pathway is not working, mutation will be colourless
 - > double mutant = colourless >> novel phenotype (new unique appearance)

Gene Interactions

- 9:7 ratio = genes in the same pathway
 - Example: harebell flowers colour



- A number of genes contribute to the phenotype
- Any gene changes in the pathway will result in the loss of pigment (pathway block)
- **Example 2: Interaction between regulatory protein and its target**
 - Regulatory protein regulates expression of another gene
 - Both WT= WT phenotype
 - Mutation in the regulatory gene= mutant
 - Mutation in the structural protein=mutant
 - Mutation in both genes=mutant
- 9:3:4 ratio = recessive epistasis
 - 2 mutants have the same phenotype
 - Double mutant shows phenotype of one but not the other
 - Epistatic = the overriding mutation
 - Hypostatic = the overridden mutation
 - Epistatic mutation is carried by gene that is farther upstream than the gene of the overridden mutation (gene in the same pathway)
- **Example: Labrador coat colour**

- **B = Black, b = brown**
- **E = normal color deposition, e = no color deposition (gold)**

P $B/B\ e/e$ (gold) \times $b/b\ E/E$ (brown) Cross btw 2 mutants

F1 $B/b\ E/e$ Black (WT)

F2 $B/-\ E/-$ 9 black (WT)
 $B/-\ e/e$ 3 gold
 $b/b\ E/-$ 3 brown
 $b/b\ e/e$ 1 gold

- Recessive phenotype (no colour deposition) overrides the colour phenotype

- 12:3:1 ratio = dominant epistasis
 - Example: flower colour

Foxgloves

- **Phenotype = petal colouration**
- **D = dark red, d = light red**
- **W = restricts colour to throat spots**

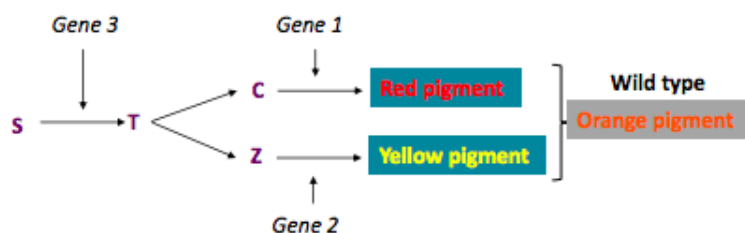
P Dark red $D/D; w/w$ \times White with spots $d/d; W/W$

F1 $D/d\ W/w$ White with spots

F2 $D/-; W/-$ 9 White with spots
 $d/d; W/-$ 3 White with spots
 $D/-; w/w$ 3 dark red
 $d/d; w/w$ 1 light red

- Dominant phenotype overrides the other phenotype

Alternative pathway: Epistasis due to divergent pathway



- 13:3 ratio
 - If recessive suppressor itself have no detectable phenotype
 - a=mutant a+=WT su+= WT su= suppressor mutation