Objectives

- Describe some violations of Mendel's laws.
- Define mutation and polymorphism, and explain their similarities and differences.
- Explain how codominance and incomplete dominance are different than Mendelian dominance.
- Compare and contrast pleiotropy and polygenic inheritance.
- Compare and contrast penetrance and expressivity.
- Define “linked” genes, and explain why genes that are located very close together on a chromosome are likely to be linked.

Non-Mendelian Genetics

Introduction

Single-gene stuff
More complex stuff

What do you mean, non-Mendelian?

Mendel was spot-on! But the traits (and genes) he happened to choose were very simple:

- Each gene had only two alleles.
- One allele was always dominant.
- Each trait had two opposing features.
- The phenotypes were always predictable.

But many genes are not so straightforward:

- Genes often have more than two alleles.
- Alleles aren’t always completely dominant or completely recessive.
- Genes don’t always act alone.
- The gene:trait relationship isn’t always 1:1.
Non-Mendelian Genetics

Introduction

Single-gene stuff

• Multiple alleles
• Codominance
• Incomplete dominance
• Pleiotropy
• Penetrance
• Expressivity

Multiple Alleles

Genes often have several common alleles. Alleles that are present in >1% of the population are called “polymorphisms.”

New alleles arise by mutation. Example: a gene for wing shape in flies

The normal (“wild-type”) allele gives the fly straight wings

A mutated (or “variant”) allele might cause the wings to be curly

“Mutation” and “polymorphism” are not interchangeable!

Mutation:
Any variation in the normal DNA sequence of a gene.
May be bad (disease-causing), silent, or good (protective).

Polymorphism:
A common, normal variation in the DNA sequence of a gene.
“Common” = present in at least 1% of the population.
“Normal” = harmless.

BUT... the distinction is not always clear-cut.
A mutation in one population may be a polymorphism in another population.
Normal Hemoglobin

- Protein sequence
- β globin subunit
- Hemoglobin molecule
- Hemoglobin molecules float around nicely

Normal Hemoglobin Sickle Hemoglobin

- Abnormal protein sequence
- Misfolded β globin subunit
- "Sticky" hemoglobin molecule
- Hemoglobin molecules polymerize, forming long crystals

BUT: patients with the sickle cell mutation have a greater resistance against malaria infection!

The sickle cell mutation is most common in parts of the world where malaria is endemic.

Non-Mendelian Genetics

Introduction

Single-gene stuff
- Multiple alleles
- Codominance

Mendelian Dominance
- One allele is always dominant; the other is always recessive.
- Heterozygotes express the phenotype of the dominant allele.

Codominance
- Both alleles are always expressed.
- Heterozygotes express the phenotypes of BOTH alleles.
Example: The ABO Blood Group System

Three alleles: A, B, and O

<table>
<thead>
<tr>
<th>A allele</th>
<th>B antigen</th>
</tr>
</thead>
<tbody>
<tr>
<td>A A</td>
<td>A antigen</td>
</tr>
<tr>
<td>B B</td>
<td>B antigen</td>
</tr>
<tr>
<td>A B</td>
<td>No antigen</td>
</tr>
</tbody>
</table>

Six possible genotypes

<table>
<thead>
<tr>
<th>A allele</th>
<th>B antigen</th>
<th>O allele</th>
</tr>
</thead>
<tbody>
<tr>
<td>A A</td>
<td>A O</td>
<td>A B</td>
</tr>
<tr>
<td>B B</td>
<td>B O</td>
<td>O O</td>
</tr>
</tbody>
</table>

Every genotype always expresses BOTH alleles!
So the ABO alleles are codominant.

Non-Mendelian Genetics

Introduction
Single-gene stuff
- Multiple alleles
- Codominance
- Incomplete dominance

Mendelian Dominance
- For a given pair of alleles, there are only two possible phenotypes.
- Heterozygotes express the phenotype of the dominant allele.

Incomplete Dominance
- For a given pair of alleles, there are three possible phenotypes!
- Heterozygotes express a phenotype that is a “blend” of the two homozygous phenotypes.
Example: Flower Color in Snapdragons
Pea flower color alleles show Mendelian dominance.
Snapdragon flower color alleles show incomplete dominance!

What about the F_2 generation?
Notice that alleles are always inherited according to Mendel’s rules (even when they show incomplete dominance).

Non-Mendelian Genetics
Introduction
Single-gene stuff
- Multiple alleles
- Codominance
- Incomplete dominance
- Pleiotropy

Pleiotropy
- Mendel’s genes each controlled a single feature.
- Pleiotropic genes affect many features.
- Note that pleiotropic alleles are still inherited in the normal, Mendelian fashion.

Mendel noticed this, actually!
Plants with purple flowers always had brown seed coats and reddish axils.
Plants with white flowers always had clear seed coats and clear axils.

Example: Marfan Syndrome
- Caused by a mutation in the fibrillin gene.
- Patients have many findings: tall height, heart valve abnormalities, aortic rupture, visual problems.
- ...so the mutated fibrillin allele is pleiotropic.
- Findings seem unrelated but actually, they make sense (fibrillin encodes a connective tissue protein).
Non-Mendelian Genetics

Introduction

Single-gene stuff
  • Multiple alleles
  • Codominance
  • Incomplete dominance
  • Pleiotropy
  • Penetrance
  • Expressivity

Penetrance and Expressivity

In Mendel’s work, plants with a particular genotype ALWAYS had the same phenotype. We now know that for some genes, this doesn’t hold true.
  • The same gene may produce different phenotypes in different people.
  • The same gene may be expressed in some people but not in others.

Penetrance

Sometimes, despite being dominant, a gene will simply not be expressed!

Complete Penetrance
  Everyone with the allele expresses the trait.

Reduced Penetrance
  Some people with the allele express the trait; some don’t.

Expressivity

Sometimes, the same genotype produces different phenotypes in different people!

Narrow Expressivity
  Same genotype, same phenotype.

Variable Expressivity
  Same genotype, different phenotypes.
Variable Expressivity in a Kindred with Marfan Syndrome

- Tall, dilated aorta
- Dislocated lenses, dilated aorta
- Tall, dislocated lenses, dilated aorta

Marfan syndrome: tall, thin patient with long limbs

Marfan syndrome: arachnodactyly

Marfan syndrome: dislocated lens

A couple physical exam signs present in Marfan syndrome:
- Thumb sticks out of fist
- Thumb and middle finger overlap when encircling wrist

Non-Mendelian Genetics

Introduction
- Single-gene stuff
- More complex stuff
  - Polygenic inheritance
  - Linkage
Non-Mendelian Genetics

Introduction

Single-gene stuff

More complex stuff

• Polygenic inheritance

Polygenic Inheritance

• In Mendel’s work, each feature (like flower color) was encoded by a single gene.
• In humans, it’s not that straightforward.
• Human features like eye color, hair color, skin color, and height are controlled by many genes (poly = many, genic = genes).
• Disease risk sometimes involves multiple genes.
• This makes things complicated, but interesting.

Eye color is not a simple Mendelian trait!

Eye color Melanocytes in the iris

It’s determined by 16 genes, most of which are involved in melanin production.

Non-Mendelian Genetics

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Single-gene stuff

More complex stuff

• Polygenic inheritance
• Linkage

Why?

Because of the random orientation of homologous chromosomes in meiosis I.

Genes on separate chromosomes assort independently, like Mendel said.
Genes far apart on the same chromosome also assort independently.

Because of crossing over ("homologous recombination").

Genes close together on the same chromosome don’t assort independently.

Because crossovers between genes that are close together are uncommon.

When genes are close together, there’s not much room for crossover.

So although recombination can happen, it’s not common.

Summary: Linkage

Unlinked genes
- Located on different chromosomes, or far apart on the same chromosome.
- Sorted into gametes independently of each other.
- Follow Mendel’s law of independent assortment.

Linked genes
- Located close to each other on the same chromosome.
- Usually sorted into gametes together as a unit.
- Violate Mendel’s law of independent assortment.