Describing Variation & Patterns of Diversity

BIOL 1435

January 31, 2023

Name, Year, Major, and what was the last song you listened to today?

Overview

1. **ATGC's of life & encoding DNA**

2. **Measures of sequence diversity**

3. **In class coding exercise**

Overview

1. ATGC's of life & encoding DNA

2. Measures of sequence diversity

3. In class coding exercise
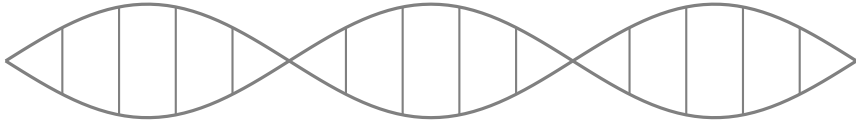
DNA consists of four nucleotides

ATGC
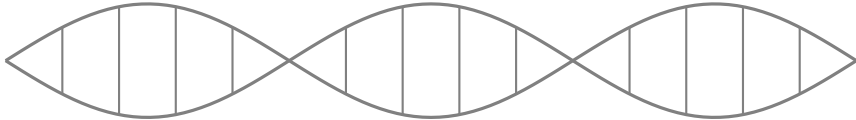
DNA is organized onto chromosomes

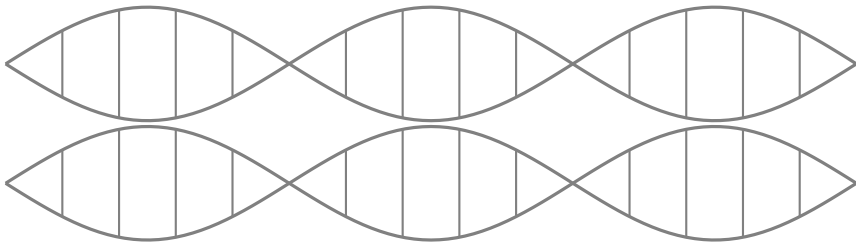# Ploidy (#N): number of sets of chromosomes

N = haploid

Ploidy (#N): number of sets of chromosomes

2N = diploid

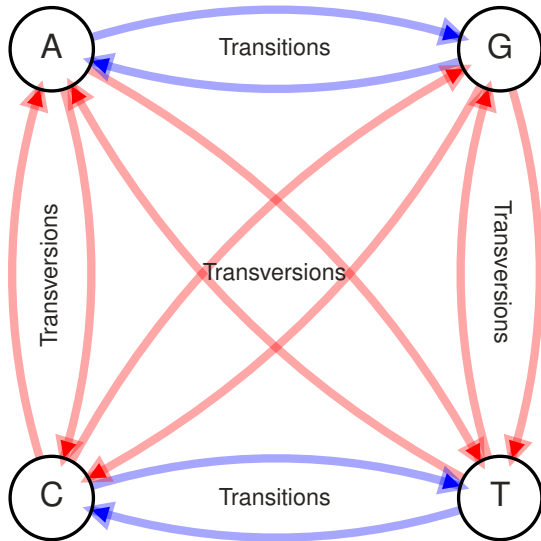# Q: How does genetic variation arise?

A: Mutations

How do we encode DNA?

$$m \text{ (sites)} \times n \text{ (chromosomes)}$$

How do we encode DNA?

$$
\begin{array}{c}
\quad ind_1 \quad ind_2 \quad ind_3 \quad ind_4 \quad ind_5 \\
\begin{array}{c} pos_1 \\ pos_2 \\ pos_3 \\ pos_4 \\ pos_5 \end{array}
\left[
\begin{array}{ccccc}
T & T & T & T & T \\
C & G & G & C & G \\
A & T & A & T & T \\
G & G & G & G & C \\
T & A & A & A & A
\end{array}
\right]
\end{array}
$$

Genotype matrices

$$
\begin{array}{c}
\\
pos_1 \\
pos_2 \\
pos_3 \\
pos_4 \\
pos_5
\end{array}
\begin{array}{ccccc}
ind_1 & ind_2 & ind_3 & ind_4 & ind_5 \\
\left[\begin{array}{ccccc}
T & T & T & T & T \\
C & G & G & C & G \\
A & T & A & T & T \\
G & G & G & G & C \\
T & A & A & A & A
\end{array}\right]
\end{array}
=
\begin{bmatrix}
0 & 0 & 0 & 0 & 0 \\
0 & 1 & 1 & 0 & 1 \\
0 & 1 & 0 & 1 & 1 \\
0 & 0 & 0 & 0 & 1 \\
1 & 0 & 0 & 0 & 0
\end{bmatrix}
$$

0 = reference or ancestral allele
1 = alternative or derived allele

Genotype matrices

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix}$$

Genotype matrices

$$
\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} = \begin{bmatrix} 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix}
$$

Some terminology...

- Single nucleotide polymorphism (SNP)
- Single nucleotide variant (SNV)
- Variant site
- Segregating site

Overview

How would you summarize this genotype matrix?

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix}$$

Measurements of genetic variation

- Segregating sites (S)
- Site frequency spectrum (SFS)
- Gene diversity (h & H)
    - Also referred to as *expected heterozygosity*
- Nucleotide diversity (Π & π)

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Segregating sites (S)

Definition
A segregating site is a site that is polymorphic in the data—i.e., there are multiple alleles observed.

Segregating sites (S)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix}$$

Segregating sites (S)

$$
\begin{bmatrix}
0 & 0 & 0 & 0 & 0 \\
0 & 1 & 1 & 0 & 1 \\
0 & 1 & 0 & 1 & 1 \\
0 & 0 & 0 & 0 & 1 \\
1 & 0 & 0 & 0 & 0
\end{bmatrix}
\rightarrow
\begin{bmatrix}
0 \\
1 \\
1 \\
1 \\
1
\end{bmatrix}
$$

Segregating sites (S)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} \rightarrow \begin{bmatrix} 0 \\ 1 \\ 1 \\ 1 \\ 1 \end{bmatrix} \rightarrow S = 4$$

Measurements of genetic variation

- Segregating sites (S)
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Site frequency spectrum (SFS)

Definition

Minor allele frequency spectrum: Histogram of the frequency of the less common allele which range from $1/n$ to 0.5 where n is the total number of chromosomes.

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Derived allele frequency spectrum: Histogram of the frequency of the derived allele—normally determined by the use of an outgroup—which range from $1/n$ to $(n-1)/n$.

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Definition

Minor allele frequency spectrum: Histogram of the frequency of the less common allele which range from $1/n$ to 0.5 where n is the total number of chromosomes.

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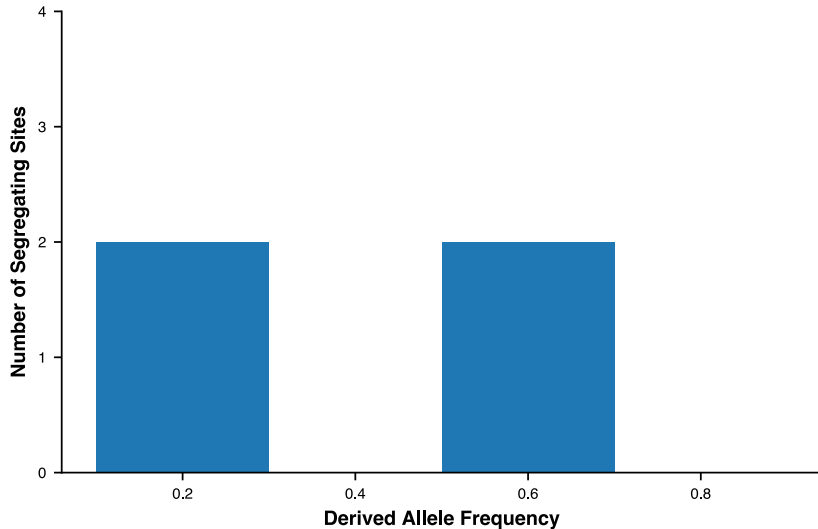
Note

Minor allele frequency spectrum = Folded SFS

Derived allele frequency spectrum = Unfolded SFS

Site frequency spectrum (SFS)

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Gene diversity (h & H)

Definition

Gene diversity is the probability that two random DNA sequences are different.

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Gene diversity is the probability that two random DNA sequences are different.

Equation

$$h = 1 - \sum_{i=1}^{m} p_i^2 \tag{1}$$

Where p_i is the frequency of the ith allele out of m observed alleles.

$$H = \frac{1}{L} \sum_{j=1}^{L} h_j \tag{2}$$

Where h_j is the gene diversity for site j and L is to the total number of sites.

Dave's tips and tricks

$$h = 1 - (p^2 + q^2)$$

Where p is the frequency of the derived/alternate allele and $q = (1 - p)$

Gene diversity (*h* & *H*)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix}$$

Gene diversity (h & H)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} \rightarrow \begin{bmatrix} 1 - \left(1^2 + 0^2\right) \\ 1 - \left(2^2/5^2 + 3^2/5^2\right) \\ 1 - \left(2^2/5^2 + 3^2/5^2\right) \\ 1 - \left(1^2/5^2 + 4^2/5^2\right) \\ 1 - \left(1^2/5^2 + 4^2/5^2\right) \end{bmatrix}$$

Gene diversity (h & H)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} \rightarrow \begin{bmatrix} 1 - \left(1^2 + 0^2\right) \\ 1 - \left(2^2/5^2 + 3^2/5^2\right) \\ 1 - \left(2^2/5^2 + 3^2/5^2\right) \\ 1 - \left(1^2/5^2 + 4^2/5^2\right) \\ 1 - \left(1^2/5^2 + 4^2/5^2\right) \end{bmatrix} \rightarrow h_j = \begin{bmatrix} 0 \\ 12/25 \\ 12/25 \\ 8/25 \\ 8/25 \end{bmatrix}$$

Gene diversity (h & H)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} \rightarrow \begin{bmatrix} 1 - \left(1^2 + 0^2\right) \\ 1 - \left(2^2/5^2 + 3^2/5^2\right) \\ 1 - \left(2^2/5^2 + 3^2/5^2\right) \\ 1 - \left(1^2/5^2 + 4^2/5^2\right) \\ 1 - \left(1^2/5^2 + 4^2/5^2\right) \end{bmatrix} \rightarrow h_j = \begin{bmatrix} 0 \\ 12/25 \\ 12/25 \\ 8/25 \\ 8/25 \end{bmatrix}$$

$$H = {}^{40}/_{25} \times {}^{1}/_{5} = {}^{8}/_{25}$$

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Definition

Nucleotide diversity is the average number of pairwise differences between genotypes drawn from the same population.

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Equation

$$\Pi = \frac{\sum_{i<j} k_{ij}}{\binom{n}{2}} \tag{3}$$

Where k_{ij} is the number of nucleotide differences between the ith and j^{th} sequence in the sample and the denominator represents the number of unique comparisons being made between n sequences.

$$\pi = \frac{\Pi}{L} \tag{4}$$

Where L is to the total number of sites.

Dave's tips and tricks

Note

$$\binom{n}{2} = \frac{n(n-1)}{2}$$

Nucleotide diversity (Π & π)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix}$$

Nucleotide diversity (Π & π)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} \rightarrow \begin{matrix} (0+0+0+0) = 0 \\ (3+1+1+1) = 6 \\ (3+1+2+0) = 6 \\ (1+1+1+1) = 4 \\ (4+0+0+0) = 4 \end{matrix}$$

Nucleotide diversity (Π & π)

$$
\begin{bmatrix}
0 & 0 & 0 & 0 & 0 \\
0 & 1 & 1 & 0 & 1 \\
0 & 1 & 0 & 1 & 1 \\
0 & 0 & 0 & 0 & 1 \\
1 & 0 & 0 & 0 & 0
\end{bmatrix}
\begin{array}{l}
(0+0+0+0) = 0 \\
(3+1+1+1) = 6 \\
\rightarrow (3+1+2+0) = 6 \rightarrow \Pi = 20 \div \frac{5(5-1)}{2} = 2 \\
(1+1+1+1) = 4 \\
(4+0+0+0) = 4
\end{array}
$$

Nucleotide diversity (Π & π)

$$\begin{bmatrix} 0 & 0 & 0 & 0 & 0 \\ 0 & 1 & 1 & 0 & 1 \\ 0 & 1 & 0 & 1 & 1 \\ 0 & 0 & 0 & 0 & 1 \\ 1 & 0 & 0 & 0 & 0 \end{bmatrix} \begin{matrix} (0+0+0+0)=0 \\ (3+1+1+1)=6 \\ \rightarrow (3+1+2+0)=6 \rightarrow \Pi = 20 \div \frac{5(5-1)}{2} = 2 \\ (1+1+1+1)=4 \\ (4+0+0+0)=4 \end{matrix}$$

$$\pi = 2 \times \text{1/5} = \text{2/5}$$

Overview