## Unit: DNA and Human Heredity (Ch. 12-14)

## "I can…"

- \_\_\_\_\_ *review* experiments leading up to the discovery of the double helix.
- \_\_\_\_\_ *create* a model that accurately depicts the structure of DNA.
- \_\_\_\_\_ *develop* a flowchart illustrating the steps of DNA replication.
- \_\_\_\_\_ explain how and why nucleotides of DNA serve as the template for mRNA (transcription).
- \_\_\_\_\_ *use* the universal genetic code to translate mRNA into a chain of amino acids (translation).
- \_\_\_\_\_ identify the types of mutations that occur in a strand of DNA.
- \_\_\_\_\_ *compile* a list of causes (intrinsic and extrinsic) of genetic mutations.
- \_\_\_\_\_ *predict* the changes in the genetic code of a strand of DNA resulting from point mutations (substitutions) and frameshift mutations (deletions, insertions, and translocations).
- \_\_\_\_\_ apply the principles of transcription and translation to normal and mutated DNA strands
- to compare and contrast the resulting sequences of amino acids.
- \_\_\_\_\_ interpret karyotypes to identify common genetic mutations
- \_\_\_\_\_ evaluate the impacts of mutations in sex cells versus body cells.

## **Essential Vocabulary/Concepts:**

adenine	rRNA (ribosomal)
anticodon	thymine
codon	transcription
cytosine	transformation
deoxyribose	translation
DNA	tRNA (transfer)
DNA polymerase	uracil
double helix	hydrogen bonds
guanine	autosomes
mRNA (messenger)	chromosomal mutation
mutation	deletion
nitrogenous base	duplication
nucleotide	frameshift mutation
phosphate	gene mutation
polypeptide chain	inversion
protein	karyotype
protein synthesis	mutation
replication	nondisjunction
ribose	point mutation
ribosome	sex chromosomes
RNA	substitution

translocation trisomy

## Key Scientists

Watson & Crick