

Identifying the Mutation

Identify the following:

- Substitution mutation:** the replacement of one nucleotide in a sequence. Substitution mutations can be categorized into three types: point mutations, silent mutations, and missense mutations. Point mutations are substitutions that change a single nucleotide, while silent mutations are substitutions that do not change the amino acid sequence. Missense mutations are substitutions that change the amino acid sequence.
- Insertion mutation:** the addition of one or more nucleotides into a sequence. Insertion mutations can be categorized into two types: point mutations and frameshift mutations. Point mutations are insertions that change a single nucleotide, while frameshift mutations are insertions that change the reading frame of the sequence.
- Deletion mutation:** the removal of one or more nucleotides from a sequence. Deletion mutations can be categorized into two types: point mutations and frameshift mutations. Point mutations are deletions that change a single nucleotide, while frameshift mutations are deletions that change the reading frame of the sequence.
- Frameshift mutation:** a mutation that shifts the reading frame of a sequence. Frameshift mutations can be caused by insertions or deletions of one or more nucleotides. Frameshift mutations can be categorized into two types: point mutations and frameshift mutations. Point mutations are frameshift mutations that change a single nucleotide, while frameshift mutations are frameshift mutations that change the reading frame of the sequence.
- Point mutation:** a mutation that changes a single nucleotide. Point mutations can be categorized into three types: silent mutations, missense mutations, and nonsense mutations. Silent mutations are point mutations that do not change the amino acid sequence. Missense mutations are point mutations that change the amino acid sequence. Nonsense mutations are point mutations that change the amino acid sequence to a stop codon.
- Frameshift mutation:** a mutation that shifts the reading frame of a sequence. Frameshift mutations can be caused by insertions or deletions of one or more nucleotides. Frameshift mutations can be categorized into two types: point mutations and frameshift mutations. Point mutations are frameshift mutations that change a single nucleotide, while frameshift mutations are frameshift mutations that change the reading frame of the sequence.

Identify the type of mutation that caused the following changes in the DNA sequence. Remember that the original DNA sequence is 5'-ATG GCG TGG-3' and the mutated DNA sequence is 5'-ATG GCG TGG-3'.

Changes in DNA Sequence	Identify the Type of Mutation
1. Original DNA: 5'-ATG GCG TGG-3' Mutated DNA: 5'-ATG GCG TGG-3'	Point mutation
2. Original DNA: 5'-ATG GCG TGG-3' Mutated DNA: 5'-ATG GCG TGG-3'	Point mutation
3. Original DNA: 5'-ATG GCG TGG-3' Mutated DNA: 5'-ATG GCG TGG-3'	Point mutation
4. Original DNA: 5'-ATG GCG TGG-3' Mutated DNA: 5'-ATG GCG TGG-3'	Point mutation
5. Original DNA: 5'-ATG GCG TGG-3' Mutated DNA: 5'-ATG GCG TGG-3'	Point mutation

Identifying Mutations Biology Unit 7 Lesson 2

Marco Cascella



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