

Identifying the Mutation

Analyze the following:

- Substitution mutation:** the change of one nucleotide in a sequence resulting in a silent mutation that has no impact on the amino acid sequence. Caused by the fact that single nucleotide mutation, at the codon level, can result in no change in the amino acid sequence or a synonymous mutation.
- Insertion mutation:** the addition of one or more nucleotides into a sequence.
- Deletion mutation:** a chromosome rearrangement that has resulted in a chromosome segment being lost.
- Translocation:** a chromosome rearrangement caused by exchange between non-homologous chromosomes.
- Transposition:** factors all types of transposable elements that can move from one location to another.
- Amplification:** a DNA sequence is duplicated many times, resulting in multiple copies.
- Point mutation:** a mutation affecting only one or two base pairs resulting in a silent mutation.
- Indel mutation:** a mutation that adds or removes one or more bases.
- Frameshift:** a genetic mutation caused by insertion or deletion of a number of nucleotides that is not a multiple of three, resulting in a shift of the reading frame.

Identify the type of mutation that caused the change in the sequence of the original and copied DNA sequences. Use the following table to help you identify the type of mutation that caused the change in the sequence of the original and copied DNA sequences.

Change in DNA Sequence	Identify the Type of Mutation
1. Original DNA: <code>ATG TAC GGC CCG</code> Copied DNA: <code>ATG TAC GGC CCG</code>	Substitution
2. Original DNA: <code>ATG TAC GGC CCG</code> Copied DNA: <code>ATG TAC GGC CCG</code>	Deletion
3. Original DNA: <code>ATG TAC GGC CCG</code> Copied DNA: <code>ATG TAC GGC CCG</code>	Insertion
4. Original DNA: <code>ATG TAC GGC CCG</code> Copied DNA: <code>ATG TAC GGC CCG</code>	Frameshift
5. Original DNA: <code>ATG TAC GGC CCG</code> Copied DNA: <code>ATG TAC GGC CCG</code>	Frameshift

Identifying Mutations Biology Unit 7 Lesson

J Elliott



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