

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 results in a loss of nucleotides, usually and in pairs.
- 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, leading to a higher copy number.
- Point mutation:** a mutation affecting only one or two base pairs resulting in a silent mutation.
- Indel mutation:** a mutation that results in a deletion or insertion of one or more nucleotides.
- Frameshift:** a genetic mutation caused by insertion or deletion of a number of nucleotides that is not a multiple of three, leading to a shift in the reading frame.

mutations, and a strategy that involves the comparison of DNA. Please use the original and mutant DNA sequences and determine whether the mutation is a substitution, insertion, deletion, or frameshift mutation. The original DNA sequence is given below.

Changes in DNA Sequence	Identify the Type of Mutation
1. Original DNA: ATG TGG CCG GCG Copied DNA: ATG TGG CCG GCG	Substitution
2. Original DNA: CGA TAC GAA GCT Copied DNA: CGA TAC GAA GCT	Point
3. Original DNA: ATG TGG CCG GCG Copied DNA: ATG TGG CCG GCG	Indel
4. Original DNA: T GAA GCT CAA GCT Copied DNA: T GAA GCT CAA GCT	Indel
5. Original DNA: C GCG CAA GCT Copied DNA: C GCG CAA GCT ATG	Frameshift

Identifying Mutations Biology Unit 7 Lesson 2 Answer

Kathleen Armour



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