

Identifying the Mutation

Identify the following:

- Substitution mutation:** the replacement of one amino acid by another without changing the sequence of amino acids. Substitutions are caused by the replacement of single nucleotides (point mutations) or the replacement of multiple nucleotides (insertions or deletions).
- Insertion mutation:** the addition of one or more nucleotides to a DNA sequence.
- Deletion mutation:** the removal of one or more nucleotides from a DNA sequence.
- Translocation:** a chromosomal rearrangement in which a segment of one chromosome is moved to a non-homologous chromosome.
- Transposition:** the movement of a segment of DNA from one location to another.
- Replication:** the process of making a copy of a DNA molecule.
- Point mutation:** a mutation that affects only one nucleotide base pair.
- Indel mutation:** a mutation that involves the insertion or deletion of one or more nucleotides.
- Frameshift:** a mutation that shifts the reading frame of a gene, resulting in a different amino acid sequence.

mutations, and a strategy that involves the comparison of the original DNA sequence with the mutated DNA sequence. The strategy involves the following steps: 1. Identify the original DNA sequence. 2. Identify the mutated DNA sequence. 3. Compare the two sequences. 4. Identify the type of mutation.

Changes in DNA Sequence	Identify the Type of Mutation
1. Original DNA: ATG CCG GGC TTT Copied DNA: ATG CCG GGC TTT	Substitution
2. Original DNA: ATG CCG GGC TTT Copied DNA: ATG CCG GGC TTT	Point
3. Original DNA: ATG CCG GGC TTT Copied DNA: ATG CCG GGC TTT	Indel
4. Original DNA: ATG CCG GGC TTT Copied DNA: ATG CCG GGC TTT	Frameshift
5. Original DNA: ATG CCG GGC TTT Copied DNA: ATG CCG GGC TTT	Frameshift

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Keith Johnson



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